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(54) Title: HUMAN ORPHAN G PROTEIN-COUPLED RECEPTORS

(57) Abstract

The invention disclosed in this patent document relates to transmembrane receptors, more particularly to endogenous, human orphan G protein-coupled receptors.

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HUMAN ORPHAN G PROTEIN-COUPLED RECEPTORS

This patent document claims priority benefit of each of the following applications, all filed with the United States Patent and Trademark Office via U.S. Express Mail on the 5 indicated filing dates: U.S. Provisional Number 60/121,852, filed; February 26, 1999 claiming the benefit of U.S. Provisional Number 60/109,213, filed November 20, 1998; U.S. Provisional Number 60/120,416, filed February 16, 1999; U.S. Provisional Number 60/123,946, filed March 12, 1999; U.S. Provisional Number 60/123,949, filed March 12, 1999; U.S. Provisional Number 60/136,436, filed May 28, 1999; U.S. Provisional 10 Number 60/136,439, filed May 28, 1999; U.S. Provisional Number 60/136,567, filed May 28, 1999; U.S. Provisional Number 60/137,127, filed May 28, 1999; U.S. Provisional Number 60/137,131, filed May 28, 1999; U.S. Provisional Number 141,448, filed June 29, 1999 claiming priority from U.S. Provisional Number 60/136,437, filed May 28, 1999; U.S. Provisional Number ___ ____ (Arena Pharmaceuticals, Inc. docket number 15 CHN10-1), filed September 29, 1999; U.S. Provisional Number 60/156,333, filed September 29, 1999; U.S. Provisional Number 60/156,555, filed September 29, 1999; U.S. Provisional Number 60/156,634, filed September 29, 1999; U.S. Provisional (Arena Pharmaceuticals, Inc. docket number RUP6-1), filed October 1, 1999; U.S. Provisional Number _____ (Arena Pharmaceuticals, Inc. docket number 20 RUP7-1), filed October 1, 1999; U.S. Provisional Number ____ (Arena Pharmaceuticals, Inc. docket number CHN6-1), filed October 1, 1999; U.S. Provisional

Number ______ (Arena Pharmaceuticals, Inc. docket number RUP5-1), filed October 1, 1999; U.S. Provisional Number ______ (Arena Pharmaceuticals, Inc. docket number CHN9-1), filed October 1, 1999. This patent document is related to U.S. Serial Number 09/170,496 filed October 13, 1998, and U.S. Serial Number unknown (Woodcock 5 Washburn Kurtz Mackiewicz & Norris, LLP docket number AREN-0054) filed on October 12, 1999 (via U.S. Express Mail) both being incorporated herein by reference. This patent document also is related to U.S. Serial No. 09/364,425; filed July 30, 1999, which is incorporated by reference in its entirety. This application also claims priority to U.S. Serial Number _____ (Woodcock, Washburn, Kurtz, Makiewicz & Norris, LLP 10 docket number AREN-0050), filed on October 12, 1999 (via U.S. Express Mail), incorporated by reference herein in its entirety. Each of the foregoing applications are incorporated herein by reference in their entirety.

FIELD OF THE INVENTION

The invention disclosed in this patent document relates to transmembrane receptors, 15 and more particularly to endogenous, orphan, human G protein-coupled receptors ("GPCRs").

BACKGROUND OF THE INVENTION

Although a number of receptor classes exist in humans, by far the most abundant and therapeutically relevant is represented by the G protein-coupled receptor (GPCR or GPCRs) 20 class. It is estimated that there are some 100,000 genes within the human genome, and of these, approximately 2% or 2,000 genes, are estimated to code for GPCRs. Receptors, including GPCRs, for which the endogenous ligand has been identified are referred to as "known" receptors, while receptors for which the endogenous ligand has not been identified

are referred to as "orphan" receptors. GPCRs represent an important area for the development of pharmaceutical products: from approximately 20 of the 100 known GPCRs, 60% of all prescription pharmaceuticals have been developed. This distinction is not merely semantic, particularly in the case of GPCRs. Thus, the orphan GPCRs are to the 5 pharmaceutical industry what gold was to California in the late 19th century – an opportunity to drive growth, expansion, enhancement and development.

GPCRs share a common structural motif. All these receptors have seven sequences of between 22 to 24 hydrophobic amino acids that form seven alpha helices, each of which spans the membrane (each span is identified by number, *i.e.*, transmembrane-1 (TM-1), 10 transmebrane-2 (TM-2), etc.). The transmembrane helices are joined by strands of amino acids between transmembrane-2 and transmembrane-3, transmembrane-4 and transmembrane-5, and transmembrane-6 and transmembrane-7 on the exterior, or "extracellular" side, of the cell membrane (these are referred to as "extracellular" regions 1, 2 and 3 (EC-1, EC-2 and EC-3), respectively). The transmembrane helices are also joined 15 by strands of amino acids between transmembrane-1 and transmembrane-2, transmembrane-3 and transmembrane-4, and transmembrane-5 and transmembrane-6 on the interior, or "intracellular" side, of the cell membrane (these are referred to as "intracellular" regions 1, 2 and 3 (IC-1, IC-2 and IC-3), respectively). The "carboxy" ("C") terminus of the receptor lies in the intracellular space within the cell, and the "amino" ("N") terminus of the receptor 20 lies in the extracellular space outside of the cell.

Generally, when an endogenous ligand binds with the receptor (often referred to as "activation" of the receptor), there is a change in the conformation of the intracellular region that allows for coupling between the intracellular region and an intracellular "G-protein." It

has been reported that GPCRs are "promiscuous" with respect to G proteins, i.e., that a GPCR can interact with more than one G protein. See, Kenakin, T., 43 Life Sciences 1095 (1988). Although other G proteins exist, currently, Gq, Gs, Gi, and Go are G proteins that have been identified. Endogenous ligand-activated GPCR coupling with the G-protein 5 begins a signaling cascade process (referred to as "signal transduction"). Under normal conditions, signal transduction ultimately results in cellular activation or cellular inhibition. It is thought that the IC-3 loop as well as the carboxy terminus of the receptor interact with the G protein.

Under physiological conditions, GPCRs exist in the cell membrane in equilibrium 10 between two different conformations: an "inactive" state and an "active" state. A receptor in an inactive state is unable to link to the intracellular signaling transduction pathway to produce a biological response. Changing the receptor conformation to the active state allows linkage to the transduction pathway (via the G-protein) and produces a biological response. A receptor may be stabilized in an active state by an endogenous ligand or a compound such 15 as a drug.

SUMMARY OF THE INVENTION

Disclosed herein are human endogenous orphan G protein-coupled receptors.

BRIEF DESCRIPTION OF THE DRAWINGS

Figures 1A and 1B provide reference "grids" for certain dot-blots provided herein 20 (see also, Figure 2A and 2B, respectively).

Figures 2A and 2B provide reproductions of the results of certain dot-blot analyses resulting from hCHN3 and hCHN8, respectively (see also, Figures 1A and 1B, respectively).

Figure 3 provides a reproduction of the results of RT-PCR analysis of hRUP3.

Figure 4 provides a reproduction of the results of RT-PCR analysis of hRUP4.

Figure 5 provides a reproduction of the results of RT-PCR analysis of hRUP6.

DETAILED DESCRIPTION

The scientific literature that has evolved around receptors has adopted a number of 5 terms to refer to ligands having various effects on receptors. For clarity and consistency, the following definitions will be used throughout this patent document. To the extent that these definitions conflict with other definitions for these terms, the following definitions shall control:

AMINO ACID ABBREVIATIONS used herein are set out in Table 1:

10		TABLE 1	
	ALANINE	ALA	A
	ARGININE	ARG	R
	ASPARAGINE	ASN	N ·
	ASPARTIC ACID	ASP	D
15	CYSTEINE	CYS	C
	GLUTAMIC ACID	GLU	E
	GLUTAMINE	GLN	$ ilde{Q}$
	GLYCINE	GLY	Ğ
	HISTIDINE	HIS	H
20	ISOLEUCINE	ILE	1
	LEUCINE	LEU	Ĺ
	LYSINE	LYS	ĸ
	METHIONINE	MET	M
	PHENYLALANINE	PHE	F
25	PROLINE	PRO	P
	SERINE	SER	S
	THREONINE	THR	r
	TRYPTOPHAN	TRP	w
	TYROSINE	TYR	v v
30	VALINE	VAL	ŷ

COMPOSITION means a material comprising at least one component.

ENDOGENOUS shall mean a material that a mammal naturally produces. ENDOGENOUS in reference to, for example and not limitation, the term "receptor," shall mean that which is naturally produced by a mammal (for example, and not limitation, a

human) or a virus. By contrast, the term **NON-ENDOGENOUS** in this context shall mean that which is not naturally produced by a mammal (for example, and not limitation, a human) or a virus.

HOST CELL shall mean a cell capable of having a Plasmid and/or Vector 5 incorporated therein. In the case of a prokaryotic Host Cell, a Plasmid is typically replicated as a autonomous molecule as the Host Cell replicates (generally, the Plasmid is thereafter isolated for introduction into a eukaryotic Host Cell); in the case of a eukaryotic Host Cell, a Plasmid is integrated into the cellular DNA of the Host Cell such that when the eukaryotic Host Cell replicates, the Plasmid replicates. Preferably, for the purposes of the invention 10 disclosed herein, the Host Cell is eukaryotic, more preferably, mammalian, and most preferably selected from the group consisting of 293, 293T and COS-7 cells.

LIGAND shall mean an endogenous, naturally occurring molecule specific for an endogenous, naturally occurring receptor.

NON-ORPHAN RECEPTOR shall mean an endogenous naturally occurring 15 molecule specific for an endogenous naturally occurring ligand wherein the binding of a ligand to a receptor activates an intracellular signaling pathway.

ORPHAN RECEPTOR shall mean an endogenous receptor for which the endogenous ligand specific for that receptor has not been identified or is not known.

PLASMID shall mean the combination of a Vector and cDNA. Generally, a Plasmid 20 is introduced into a Host Cell for the purposes of replication and/or expression of the cDNA as a protein.

VECTOR sin reference to cDNA shall mean a circular DNA capable of incorporating at least one cDNA and capable of incorporation into a Host Cell.

The order of the following sections is set forth for presentational efficiency and is not intended, nor should be construed, as a limitation on the disclosure or the claims to follow.

Identification of Human GPCRs

The efforts of the Human Genome project have led to the identification of a plethora of information regarding nucleic acid sequences located within the human genome; it has been the case in this endeavor that genetic sequence information has been made available without an understanding or recognition as to whether or not any particular genomic sequence does or may contain open-reading frame information that translate human proteins.

10 Several methods of identifying nucleic acid sequences within the human genome are within the purview of those having ordinary skill in the art. For example, and not limitation, a variety of GPCRs, disclosed herein, were discovered by reviewing the GenBankTM database, while other GPCRs were discovered by utilizing a nucleic acid sequence of a GPCR, previously sequenced, to conduct a BLASTTM search of the EST database. Table A, below, 15 lists the disclosed endogenous orphan GPCRs along with a GPCR's respective homologous GPCR:

TABLE A Disclosed Accession Open Reading Per Cent Reference To Human Number Frame Homology Homologous : 20 Orphan Identified (Base Pairs) To Designated GPCR-**GPCRs GPCR** (Accession No.) hARE-3 AL033379 1,260 bp 52.3% LPA-R U92642 hARE-4 AC006087 1,119 bp AF000546 36% P2Y5

			-8-		
4 .	hARE-5	AC006255	1,104 bp	32% Oryzias	D43633
•				latipes	
	hGPR27	AA775870	1,128 bp	var.pes	•
	hARE-1	AI090920	999 bp	43%	D13626
				42,70	D13020
				KIAA0001	•
	hARE-2	AA359504	1,122 bp	53% GPR27	
5	hPPR1	H67224	1,053 bp	39% EBI1	F 2 1 5 0 1
	hG2A	AA754702	1,113 bp	31% GPR4	L31581
	hRUP3	AL035423	1,005 bp		L36148
	ARCO15	AD000420	1,005 bp	30%	2133653
				Drosophila	
					•
	hRUP4	A 1207/50	1.0061	melanogaster	
	RKUF4	AI307658	1,296 bp	32% pNPGPR	NP_004876
			•	28% and 29 %	AAC41276
	•			Zebra fish Ya	
				zeora jun 1a	and
				and Yb.	
			٠		AAB94616
		• •		respectively	
	hRUP5	AC005849	1,413 bp	25% DEZ	Q99788
		• •			
				23% FMLPR	P21462
10	hRUP6	AC005871	1,245 bp	48% GPR66	NP_006047
	hRUP7	AC007922	1,173 bp	43% H3R	AF140538
•	hCHN3	EST 36581	1,113 bp	53% GPR27	•
	hCHN4	AA804531	1,077 bp	32% thrombin	4503637
	hCHN6	EST 2134670	1,503 bp	36% edg-1	NP 001391
15	hCHN8	EST 764455	1,029 bp	47%	D13626
			•	KIAA0001	
	hCHN9	EST 1541536	1,077 bp	41% LTB4R	NM_000752
	hCHN10	EST 1365839	1,055 bp	35% P2Y	NM 002563
• 1.		•	₹ .		

Receptor homology is useful in terms of gaining an appreciation of a role of the disclosed receptors within the human body. Additionally, such homology can provide insight 20 as to possible endogenous ligand(s) that may be natural activators for the disclosed orphan GPCRs.

B. Receptor Screening

Techniques have become more readily available over the past few years for

endogenous-ligand identification (this, primarily, for the purpose of providing a means of conducting receptor-binding assays that require a receptor's endogenous ligand) because the traditional study of receptors has always proceeded from the a priori assumption (historically based) that the endogenous ligand must first be identified before discovery could proceed to 5 find antagonists and other molecules that could affect the receptor. Even in cases where an antagonist might have been known first, the search immediately extended to looking for the endogenous ligand. This mode of thinking has persisted in receptor research even after the discovery of constitutively activated receptors. What has not been heretofore recognized is that it is the active state of the receptor that is most useful for discovering agonists, partial 10 agonists, and inverse agonists of the receptor. For those diseases which result from an overly active receptor or an under-active receptor, what is desired in a therapeutic drug is a compound which acts to diminish the active state of a receptor or enhance the activity of the receptor, respectively, not necessarily a drug which is an antagonist to the endogenous ligand. This is because a compound that reduces or enhances the activity of the active receptor state 15 need not bind at the same site as the endogenous ligand. Thus, as taught by a method of this invention, any search for therapeutic compounds should start by screening compounds against the ligand-independent active state.

As is known in the art, GPCRs can be "active" in their endogenous state even without the binding of the receptor's endogenous ligand thereto. Such naturally-active receptors can 20 be screened for the direct identification (*i.e.*, without the need for the receptor's endogenous ligand) of, in particular, inverse agonists. Alternatively, the receptor can be "activated" via, *e.g.*, mutation of the receptor to establish a non-endogenous version of the receptor that is active in the absence of the receptor's endogenous ligand.

Screening candidate compounds against an endogenous or non-endogenous, constitutively activated version of the human orphan GPCRs disclosed herein can provide for the direct identification of candidate compounds which act at this cell surface receptor, without requiring use of the receptor's endogenous ligand. By determining areas within 5 the body where the endogenous version of human GPCRs disclosed herein is expressed and/or over-expressed, it is possible to determine related disease/disorder states which are associated with the expression and/or over-expression of the receptor; such an approach is disclosed in this patent document.

With respect to creation of a mutation that may evidence constitutive activation of 10 human orphan GPCRs disclosed herein is based upon the distance from the proline residue at which is presumed to be located within TM6 of the GPCR typically nears the TM6/IC3 interface (such proline residue appears to be quite conserved). By mutating the amino acid residue located 16 amino acid residues from this residue (presumably located in the IC3 region of the receptor) to, most preferably, a lysine residue, such activation may be obtained. 15 Other amino acid residues may be useful in the mutation at this position to achieve this objective.

C. Disease/Disorder Identification and/or Selection

Preferably, the DNA sequence of the human orphan GPCR can be used to make a probe for (a) dot-blot analysis against tissue-mRNA, and/or (b) RT-PCR identification of 20 the expression of the receptor in tissue samples. The presence of a receptor in a tissue source, or a diseased tissue, or the presence of the receptor at elevated concentrations in diseased tissue compared to a normal tissue, can be preferably utilized to identify a correlation with a treatment regimen, including but not limited to, a disease associated

with that disease. Receptors can equally well be localized to regions of organs by this technique. Based on the known functions of the specific tissues to which the receptor is localized, the putative functional role of the receptor can be deduced.

D. Screening of Candidate Compounds

1. Generic GPCR screening assay techniques

When a G protein receptor becomes constitutively active (i.e., active in the absence of endogenous ligand binding thereto), it binds to a G protein (e.g., Gq, Gs, Gi, Go) and stimulates the binding of GTP to the G protein. The G protein then acts as a GTPase and slowly hydrolyzes the GTP to GDP, whereby the receptor, under normal conditions, becomes 10 deactivated. However, constitutively activated receptors continue to exchange GDP to GTP. A non-hydrolyzable analog of GTP, [35S]GTPγS, can be used to monitor enhanced binding to membranes which express constitutively activated receptors. It is reported that [35S]GTPγS can be used to monitor G protein coupling to membranes in the absence and presence of ligand. An example of this monitoring, among other examples well-known and 15 available to those in the art, was reported by Traynor and Nahorski in 1995. The preferred use of this assay system is for initial screening of candidate compounds because the system is generically applicable to all G protein-coupled receptors regardless of the particular G protein that interacts with the intracellular domain of the receptor.

2. Specific GPCR screening assay techniques

Once candidate compounds are identified using the "generic" G protein-coupled receptor assay (i.e., an assay to select compounds that are agonists, partial agonists, or inverse agonists), further screening to confirm that the compounds have interacted at the receptor site is preferred. For example, a compound identified by the "generic" assay may not bind to the

receptor, but may instead merely "uncouple" the G protein from the intracellular domain.

a. Gs and Gi.

Gs stimulates the enzyme adenylyl cyclase. Gi (and Go), on the other hand, inhibit Adenylyl cyclase catalyzes the conversion of ATP to cAMP; thus, 5 constitutively activated GPCRs that couple the Gs protein are associated with increased cellular levels of cAMP. On the other hand, constitutively activated GPCRs that couple the Gi (or Go) protein are associated with decreased cellular levels of cAMP. See, generally, "Indirect Mechanisms of Synaptic Transmission," Chpt. 8, From Neuron To Brain (3rd Ed.) Nichols, J.G. et al eds. Sinauer Associates, Inc. (1992). Thus, assays that detect cAMP can 10 be utilized to determine if a candidate compound is, e.g., an inverse agonist to the receptor (i.e., such a compound would decrease the levels of cAMP). A variety of approaches known in the art for measuring cAMP can be utilized; a most preferred approach relies upon the use of anti-cAMP antibodies in an ELISA-based format. Another type of assay that can be utilized is a whole cell second messenger reporter system assay. Promoters on genes drive 15 the expression of the proteins that a particular gene encodes. Cyclic AMP drives gene expression by promoting the binding of a cAMP-responsive DNA binding protein or transcription factor (CREB) which then binds to the promoter at specific sites called cAMP response elements and drives the expression of the gene. Reporter systems can be constructed which have a promoter containing multiple cAMP response elements before the reporter 20 gene, e.g., β -galactosidase or luciferase. Thus, a constitutively activated Gs-linked receptor causes the accumulation of cAMP that then activates the gene and expression of the reporter protein. The reporter protein such as β-galactosidase or luciferase can then be detected using standard biochemical assays (Chen et al. 1995).

Go and Gq.

Gq and Go are associated with activation of the enzyme phospholipase C, which in turn hydrolyzes the phospholipid PIP₂, releasing two intracellular messengers: 5 diacycloglycerol (DAG) and inistol 1,4,5-triphoisphate (IP₃). Increased accumulation of IP₃ is associated with activation of Gq- and Go-associated receptors. See, generally, "Indirect Mechanisms of Synaptic Transmission," Chpt. 8, From Neuron To Brain (3rd Ed.) Nichols, J.G. et al eds. Sinauer Associates, Inc. (1992). Assays that detect IP₃ accumulation can be utilized to determine if a candidate compound is, e.g., an inverse agonist to a Gq- or Go-10 associated receptor (i.e., such a compound would decrease the levels of IP₃). Gq-associated receptors can also been examined using an AP1 reporter assay in that Gq-dependent phospholipase C causes activation of genes containing AP1 elements; thus, activated Gq-associated receptors will evidence an increase in the expression of such genes, whereby inverse agonists thereto will evidence a decrease in such expression, and agonists will 15 evidence an increase in such expression. Commercially available assays for such detection are available.

3. GPCR Fusion Protein

The use of an endogenous, constitutively activated orphan GPCR, or a non-endogenous, constitutively activated orphan GPCR, for screening of candidate compounds 20 for the direct identification of inverse agonists, agonists and partial agonists provides a unique challenge in that, by definition, the receptor is active even in the absence of an endogenous ligand bound thereto. Thus, it is often useful that an approach be utilized that can enhance the signal obtained by the activated receptor. A preferred approach is the use of a GPCR Fusion Protein.

Generally, once it is determined that a GPCR is or has been constitutively activated, using the assay techniques set forth above (as well as others), it is possible to determine the predominant G protein that couples with the endogenous GPCR. Coupling of the G protein to the GPCR provides a signaling pathway that can be assessed. Because it is most preferred 5 that screening take place by use of a mammalian expression system, such a system will be expected to have endogenous G protein therein. Thus, by definition, in such a system, the constitutively activated orphan GPCR will continuously signal. In this regard, it is preferred that this signal be enhanced such that in the presence of, e.g., an inverse agonist to the receptor, it is more likely that it will be able to more readily differentiate, particularly in the 10 context of screening, between the receptor when it is contacted with the inverse agonist.

The GPCR Fusion Protein is intended to enhance the efficacy of G protein coupling with the GPCR. The GPCR Fusion Protein is preferred for screening with a non-endogenous, constitutively activated GPCR because such an approach increases the signal that is most preferably utilized in such screening techniques, although the GPCR Fusion 15 Protein can also be (and preferably is) used with an endogenous, constitutively activated GPCR. This is important in facilitating a significant "signal to noise" ratio; such a significant ratio is import preferred for the screening of candidate compounds as disclosed herein.

The construction of a construct useful for expression of a GPCR Fusion Protein is within the purview of those having ordinary skill in the art. Commercially available 20 expression vectors and systems offer a variety of approaches that can fit the particular needs of an investigator. The criteria of importance for such a GPCR Fusion Protein construct is that the GPCR sequence and the G protein sequence both be in-frame (preferably, the sequence for the GPCR is upstream of the G protein sequence) and that the "stop" codon of

the GPCR must be deleted or replaced such that upon expression of the GPCR, the G protein can also be expressed. The GPCR can be linked directly to the G protein, or there can be spacer residues between the two (preferably, no more than about 12, although this number can be readily ascertained by one of ordinary skill in the art). We have a preference (based 5 upon convenience) of use of a spacer in that some restriction sites that are not used will, effectively, upon expression, become a spacer. Most preferably, the G protein that couples to the GPCR will have been identified prior to the creation of the GPCR Fusion Protein construct. Because there are only a few G proteins that have been identified, it is preferred that a construct comprising the sequence of the G protein (i.e., a universal G protein 10 construct) be available for insertion of an endogenous GPCR sequence therein; this provides for efficiency in the context of large-scale screening of a variety of different endogenous GPCRs having different sequences.

E. Other Utility

Although a preferred use of the human orphan GPCRs disclosed herein may be for 15 the direct identification of candidate compounds as inverse agonists, agonists or partial agonists (preferably for use as pharmaceutical agents), these versions of human GPCRs can also be utilized in research settings. For example, *in vitro* and *in vivo* systems incorporating GPCRs can be utilized to further elucidate and understand the roles these receptors play in the human condition, both normal and diseased, as well as understanding the role of 20 constitutive activation as it applies to understanding the signaling cascade. The value in human orphan GPCRs is that its utility as a research tool is enhanced in that by determining the location(s) of such receptors within the body, the GPCRs can be used to understand the role of these receptors in the human body before the endogenous ligand therefor is identified.

Other uses of the disclosed receptors will become apparent to those in the art based upon, inter alia, a review of this patent document.

EXAMPLES

The following examples are presented for purposes of elucidation, and not limitation, 5 of the present invention. While specific nucleic acid and amino acid sequences are disclosed herein, those of ordinary skill in the art are credited with the ability to make minor modifications to these sequences while achieving the same or substantially similar results reported below. Unless otherwise indicated below, all nucleic acid sequences for the disclosed endogenous orphan human GPCRs have been sequenced and verified. For 10 purposes of equivalent receptors, those of ordinary skill in the art will readily appreciate that conservative substitutions can be made to the disclosed sequences to obtain a functionally equivalent receptor.

Example 1 ENDOGENOUS HUMAN GPCRS

1. Identification of Human GPCRs

Several of the disclosed endogenous human GPCRs were identified based upon a review of the GenBank database information. While searching the database, the following cDNA clones were identified as evidenced below.

	Disclosed	Accession	Complete DNA	Open Reading	Nucleic Acid	Amino
20	Human	Number	Sequence	Frame	SEQ.ID.	Acid
	Orphan		(Base Pairs)	(Base Pairs)	NO.	SEQ.ID.
	GPCRs		÷ .	:	8 m 2 m 1 m	NO.

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	hARE-3	AL033379	111,389 bp	1,260 bp	1	. 2
	hARE-4	AC006087	226,925 bp	1,119 bp	3	4
	hARE-5	AC006255	127,605 bp	1,104 bp	5 •	6
	hRUP3	AL035423	140,094 bp	1,005 bp	7	8
5	hRUP5	AC005849	169,144 bp	1,413 bp	9	10
•	hRUP6	AC005871	218,807 bp	1,245 bp	11	12
	hRUP7	AC007922	158,858 bp	1,173 bp	13	14

Other disclosed endogenous human GPCRs were identified by conducting a BLAST search of EST database (dbest) using the following EST clones as query sequences. The 10 following EST clones identified were then used as a probe to screen a human genomic library.

	Disclosed	Query	EST Clone/	Open	Nucleic Acid	Amino Acid
	Human	(Sequence)	Accession No.	Reading	SEQ.ID.NO.	SEQ.ID.NO.
	Orphan		Identified	Frame		
15	GPCRs hGPCR27	Mouse	AA775870	(Base Pairs) 1,125 bp	. 73. 2 15	16
	hARE-1	GPCR27 TDAG	1689643	999 bp	17	18
	hARE-2	GPCR27	A1090920 68530	1,122 bp	19	20
	hPPR1	Bovine	AA359504 238667	1,053 bp	21	22
20	hG2A	PPR1 Mouse	H67224 See Example 2(a),	1,113 bp	23	24
		1179426	below			

2. Full Length Cloning

N.A. = "not applicable".

a. hG2A (Seq. Id. Nos. 23 & 24)

Mouse EST clone 1179426 was used to obtain a human genomic clone containing all but three amino acid hG2A coding sequences. The 5'end of this coding sequence was obtained by using 5'RACE™, and the template for PCR was Clontech's Human Spleen Marathon-ready™ cDNA. The disclosed human G2A was amplified by PCR using the G2A 15 cDNA specific primers for the first and second round PCR as shown in SEQ.ID.NO.: 39 and SEQ.ID.NO.:40 as follows:

5'-CTGTGTACAGCAGTTCGCAGAGTG-3' (SEQ.ID.NO.: 39; 1st round PCR)

5'-GAGTGCCAGGCAGAGCAGGTAGAC-3' (SEQ.ID.NO.: 40; second round PCR).

PCR was performed using Advantage™ GC Polymerase Kit (Clontech; manufacturing 20 instructions will be followed), at 94°C for 30 sec followed by 5 cycles of 94°C for 5 sec and 72°C for 4 min; and 30 cycles of 94° for 5 sec and 70° for 4 min. An approximate 1.3 Kb PCR fragment was purified from agarose gel, digested with Hind III and Xba I and cloned into the expression vector pRC/CMV2 (Invitrogen). The cloned-insert was sequenced using the T7 Sequenase™ kit (USB Amersham; manufacturer instructions will be followed) and

the sequence was compared with the presented sequence. Expression of the human G2A will be detected by probing an RNA dot blot (Clontech; manufacturer instructions will be followed) with the P³²-labeled fragment.

b. hCHN9 (Seq. Id. Nos. 33 & 34)

Sequencing of the EST clone 1541536 indicated that hCHN9 is a partial cDNA clone having only an initiation codon; *i.e.*, the termination codon was missing. When hCHN9 was used to "blast" against the data base (nr), the 3' sequence of hCHN9 was 100% homologous to the 5' untranslated region of the leukotriene B4 receptor cDNA, which contained a termination codon in the frame with hCHN9 coding sequence. To 10 determine whether the 5' untranslated region of LTB4R cDNA was the 3' sequence of hCHN9, PCR was performed using primers based upon the 5' sequence flanking the initiation codon found in hCHN9 and the 3' sequence around the termination codon found in the LTB4R 5' untranslated region. The 5' primer sequence utilized was as follows: 5'-CCCGAATTCCTGCTTGCTCCCAGCTTGGCCC-3' (SEQ.ID.NO.: 41; sense) and 15 5'-TGTGGATCCTGCTGCTCAAAGGTCCCATTCCGG-3' (SEQ.ID.NO.: 42; antisense).

PCR was performed using thymus cDNA as a template and rTth polymerase (Perkin Elmer) with the buffer system provided by the manufacturer, 0.25 uM of each primer, and 0.2 mM of each 4 nucleotides. The cycle condition was 30 cycles of 94°C for 1 min, 65°C for 1 min and 72 °C for 1 min and 10 sec. A 1.1kb fragment consistent with the predicted size was

20 obtained from PCR. This PCR fragment was subcloned into pCMV (see below) and sequenced (see, SEQ.ID.NO.: 33).

c. hRUP 4 (Seq. Id. Nos. 37 & 38)

The full length hRUP4 was cloned by RT-PCR with human brain cDNA (Clontech)

as templates:

5'-TCACAATGCTAGGTGTGGTC-3' (SEQ.ID.NO.: 43; sense) and

5'-TGCATAGACAATGGGATTACAG-3' (SEQ.ID.NO.: 44; antisense).

PCR was performed using TaqPlus™ Precision™ polymerase (Stratagene; manufacturing 5 instructions will be followed) by the following cycles: 94°C for 2 min; 94°C 30 sec; 55°C for 30 sec, 72°C for 45 sec, and 72°C for 10 min. Cycles 2 through 4 were repeated 30 times.

The PCR products were separated on a 1% agarose gel and a 500 bp PCR fragment was isolated and cloned into the pCRII-TOPO vector (Invitrogen) and sequenced using the 10 T7 DNA Sequenase™ kit (Amsham) and the SP6/T7 primers (Stratagene). Sequence analysis revealed that the PCR fragment was indeed an alternatively spliced form of AI307658 having a continuous open reading frame with similarity to other GPCRs. The completed sequence of this PCR fragment was as follows:

5'-TCACAATGCTAGGTGTGGTCTGGCTGGTGGCAGTCATCGTAGGATCACCCATGTGGCAC
15 GTGCAACAACTTGAGATCAAATATGACTTCCTATATGAAAAGGAACACATCTGCTGCTTAGAA
GAGTGGACCAGCCCTGTGCACCAGAAGATCTACACCACCTTCATCCTTGTCATCCTCTCC
TGCCTCTTATGGTGATGCTTATTCTGTACGTAAAATTGGTTATGAACTTTGGATAAAGAAAAGA
GTTGGGGATGGTTCAGTGCTTCGAACTATTCATGGAAAAAGAAATGTCCAAAATAGCCAGGAAG
AAGAAACGAGCTGTCATTATGATGGTGACAGTGGTGGCTCTCTTTGCTGTGTGCTGGGCACCA
20 TTCCATGTTGTCCATATGATGATTGAATACAGTAATTTTGAAAAGGAATATGATGATGTCACA
ATCAAGATGATTTTTGCTATCGTGCAAATTATTGGATTTTCCAACTCCATCTGTAATCCCATTG
TCTATGCA-3' (SEQ.ID.NO.: 45)

Based on the above sequence, two sense oligonucleotide primer sets:

5'-CTGCTTAGAAGAGTGGACCAG-3' (SEQ.ID.NO.: 46; oligo 1),

25 5'-CTGTGCACCAGAAGATCTACAC-3' (SEQ.IDNO.: 47; oligo 2)

and two antisense oligonucleotide primer sets:

5'-CAAGGATGAAGGTGGTGTAGA-3' (SEQ.ID.NO.: 48; oligo 3)

5'-GTGTAGATCTTCTGGTGCACAGG-3' (SEQ.ID.NO.: 49; oligo 4)

were used for 3'- and 5'-race PCR with a human brain Marathon-Ready™ cDNA (Clontech,

Cat# 7400-1) as template, according to manufacture's instructions. DNA fragments generated by the RACE PCR were cloned into the pCRII-TOPO™ vector (Invitrogen) and sequenced using the SP6/T7 primers (Stratagene) and some internal primers. The 3' RACE product contained a poly(A) tail and a completed open reading frame ending at a TAA stop 5 codon. The 5' RACE product contained an incomplete 5' end; *i.e.*, the ATG initiation codon was not present.

Based on the new 5' sequence, oligo 3 and the following primer:

5'-GCAATGCAGGTCATAGTGAGC -3' (SEQ.ID.NO.: 50; oligo 5)

were used for the second round of 5' RACE PCR and the PCR products were analyzed as

10 above. A third round of 5' RACE PCR was carried out utilizing antisense primers:

5'-TGGAGCATGGTGACGGGAATGCAGAAG-3' (SEQ.ID.NO.: 51; oligo 6) and

5'-GTGATGAGCAGGTCACTGAGCGCCAAG-3' (SEQ.ID.NO.: 52; oligo7).

The sequence of the 5' RACE PCR products revealed the presence of the initiation codon

ATG, and further round of 5' RACE PCR did not generate any more 5' sequence. The

15 completed 5' sequence was confirmed by RT-PCR using sense primer

5'-GCAATGCAGGCGCTTAACATTAC-3' (SEQ.ID.NO.: 53; oligo 8)

and oligo 4 as primers and sequence analysis of the 650 bp PCR product generated from human brain and heart cDNA templates (Clontech, Cat# 7404-1). The completed 3' sequence was confirmed by RT-PCR using oligo 2 and the following antisense primer:

205'-TTGGGTTACAATCTGAAGGGCA-3' (SEQ.ID.NO.: 54; oligo 9)

and sequence analysis of the 670 bp PCR product generated from human brain and heart cDNA templates. (Clontech, Cat# 7404-1).

d. hRUP5 (Seq. Id. Nos. 9 & 10)

The full length hRUP5 was cloned by RT-PCR using a sense primer upstream from

ATG, the initiation codon (SEQ.ID.NO.: 55), and an antisense primer containing TCA as the stop codon (SEQ.ID.NO.: 56), which had the following sequences:

- 5'-ACTCCGTGTCCAGCAGGACTCTG-3' (SEQ.ID.NO.:55)
- 5'-TGCGTGTTCCTGGACCCTCACGTG-3' (SEQ.ID.NO.: 56)
- 5 and human peripheral leukocyte cDNA (Clontech) as a template. Advantage cDNA polymerase (Clontech) was used for the amplification in a 50ul reaction by the following cycle with step 2 through step 4 repeated 30 times: 94°C for 30 sec; 94° for 15 sec; 69° for 40 sec; 72°C for 3 min; and 72°C fro 6 min. A 1.4kb PCR fragment was isolated and cloned with the pCRII-TOPO™ vector (Invitrogen) and completely sequenced using the T7 DNA 10 Sequenase™ kit (Amsham). See, SEQ.ID.NO.: 9.

e. hRUP6 (Seq. Id. Nos. 11 & 12)

The full length hRUP6 was cloned by RT-PCR using primers:

- 5'-CAGGCCTTGGATTTTAATGTCAGGGATGG-3' (SEQ.ID.NO.: 57) and
- 5'-GGAGAGTCAGCTCTGAAAGAATTCAGG-3' (SEQ.ID.NO.: 58);
- 15 and human thymus Marathon-Ready™ cDNA (Clontech) as a template. Advantage cDNA polymerase (Clontech, according to manufacturer's instructions) was used for the amplification in a 50ul reaction by the following cycle: 94°C for 30sec; 94°C for 5 sec; 66°C for 40sec; 72°C for 2.5 sec and 72°C for 7 min. Cycles 2 through 4 were repeated 30 times. A 1.3 Kb PCR fragment was isolated and cloned into the pCRII-TOPO™ vector (Invitrogen) 20 and completely sequenced (see, SEQ.ID.NO.: 11) using the ABI Big Dye Terminator™ kit (P.E. Biosystem).

f. hRUP7 (Seq. Id. Nos. 13 & 14)

The full length RUP7 was cloned by RT-PCR using primers: 5'-TGATGTGATGCCAGATACTAATAGCAC-3' (SEQ.ID.NO.: 59; sense) and

5'-CCTGATTCATTTAGGTGAGATTGAGAC-3' (SEQ.ID.NO.: 60; antisense)
and human peripheral leukocyte cDNA (Clontech) as a template. Advantage™ cDNA
polymerase (Clontech) was used for the amplification in a 50 ul reaction by the following
cycle with step 2 to step 4 repeated 30 times: 94°C for 2 minutes; 94°C for 15 seconds; 60°C
5 for 20 seconds; 72°C for 2 minutes; 72°C for 10 minutes. A 1.25 Kb PCR fragment was
isolated and cloned into the pCRII-TOPO™ vector (Invitrogen) and completely sequenced
using the ABI Big Dye Terminator™ kit (P.E. Biosystem). See, SEQ.ID.NO.: 13.

g. hARE-5 (Seq. Id. Nos. 5 & 6)

The full length hARE-5 was cloned by PCR using the hARE5 specific primers 10 5'-CAGCGCAGGGTGAAGCCTGAGAGC-3' SEQ.ID.NO.: 69 (sense, 5' of initiation codon ATG) and 5'-GGCACCTGCTGTGCAGG-3' SEQ.ID.NO.:70 (antisense, 3' of stop codon TGA) and human genomic DNA as template. TaqPlus Precision™ DNA polymerase (Stratagene) was used for the amplification by the following cycle with step 2 to step 4 repeated 35 times: 96°C, 2 minutes; 96°C, 20 seconds; 58°C, 30 seconds; 72°C, 2 minutes; and 72°C, 10 minutes

A 1.1 Kb PCR fragment of predicated size was isolated and cloned into the pCRII-TOPO™ vector (Invitrogen) and completely sequenced (SEQ.ID.NO.:5) using the T7 DNA Sequenase™ kit (Amsham).

h. hARE-4 (Seq. Id. Nos.: 3 & 4)

The full length hARE-4 was cloned by PCR using the hARE-4 specific primers 5'-20 CTGGTGTCCCATGGCATCCC-3' SEQ.ID.NO.:67 (sense, 5' of initiation codon ATG) and 5'-GTAAGCCTCCCAGAACGAGAGG-3' SEQ.ID.NO.: 68 (antisense, 3' of stop codon TGA) and human genomic DNA as template. Taq DNA polymerase (Stratagene) and 5% DMSO was used for the amplification by the following cycle with step 2 to step 3 repeated 35 times:

94°C, 3 minutes; 94°C, 30 seconds; 59°C, 2 minutes; 72°C, 10 minutes

A 1.12 Kb PCR fragment of predicated size was isolated and cloned into the pCRII-TOPOTM vector (Invitrogen) and completely sequenced (SEQ.ID.NO.:3) using the T7 DNA SequenaseTM kit (Amsham).

i. hARE-3 (Seq.Id.Nos.: 1 & 2)

The full length hARE-3 was cloned by PCR using the hARE-3 specific primers 5'-gatcaagettCCATCCTACTGAAACCATGGTC-3' SEQ.ID.NO.:65 (sense, lower case nucleotides represent Hind III overhang, ATG as initiation codon) and 5'-gatcagatctCAGTTCCAATATTCACACCACCGTC-3' SEQ.ID.NO.:66 (antisense, lower case 10 nucleotides represent Xba I overhang, TCA as stop codon) and human genomic DNA as template. TaqPlus PrecisionTM DNA polymerase (Stratagene) was used for the amplification by the following cycle with step 2 to step 4 repeated 35 times: 94°C, 3 minutes; 94°C, 1 minute; 55°C, 1 minute; 72°C, 2 minutes; 72°C, 10 minutes.

A 1.3 Kb PCR fragment of predicated size was isolated and digested with Hind III 15 and Xba I, cloned into the pRC/CMV2 vector (Invitrogen) at the Hind III and Xba I sites and completely sequenced (SEQ.ID.NO.:1) using the T7 DNA SequenaseTM kit (Amsham).

j. hRUP3 (Seq. Id. Nos.: 7 & 8)

The full length hRUP3 was cloned by PCR using the hRUP3 specific primers 5'-GTCCTGCCACTTCGAGACATGG-3' SEQ.ID.NO.:71 (sense, ATG as initiation codon) and 5'-20 GAAACTTCTCTGCCCTTACCGTC-3' SEQ.ID.NO.:72 (antisense, 3' of stop codon TAA) and human genomic DNA as template. TaqPlus Precision™ DNA polymerase (Stratagene) was used for the amplification by the following cycle with step 2 to step 4 repeated 35 times: 94°C, 3 minutes; 94°C, 1 minute; 58°C, 1 minute; 72°C, 2 minutes; 72°C, 10 minutes

A 1.0 Kb PCR fragment of predicated size was isolated and cloned into the pCRII-TOPO™ vector (Invitrogen) and completely sequenced (SEQ.ID.NO.: 7)using the T7 DNA sequenase kit (Amsham).

Example 2 5 RECEPTOR EXPRESSION

Although a variety of cells are available to the art for the expression of proteins, it is most preferred that mammalian cells be utilized. The primary reason for this is predicated upon practicalities, *i.e.*, utilization of, *e.g.*, yeast cells for the expression of a GPCR, while possible, introduces into the protocol a non-mammalian cell which may not (indeed, in the 10 case of yeast, does not) include the receptor-coupling, genetic-mechanism and secretary pathways that have evolved for mammalian systems – thus, results obtained in non-mammalian cells, while of potential use, are not as preferred as that obtained from mammalian cells. Of the mammalian cells, COS-7, 293 and 293T cells are particularly preferred, although the specific mammalian cell utilized can be predicated upon the particular 15 needs of the artisan. The general procedure for expression of the disclosed GPCRs is as follows.

On day one, 1X10⁷ 293T cells per 150mm plate were plated out. On day two, two reaction tubes will be prepared (the proportions to follow for each tube are per plate): tube A will be prepared by mixing 20µg DNA (e.g., pCMV vector; pCMV vector with receptor 20 cDNA, etc.) in 1.2ml serum free DMEM (Irvine Scientific, Irvine, CA); tube B will be prepared by mixing 120µl lipofectamine (Gibco BRL) in 1.2ml serum free DMEM. Tubes A and B are admixed by inversions (several times), followed by incubation at room temperature for 30-45min. The admixture can be referred to as the "transfection mixture". Plated 293T cells are washed with 1XPBS, followed by addition of 10ml serum free DMEM.

2.4ml of the transfection mixture will then be added to the cells, followed by incubation for 4hrs at $37^{\circ}\text{C}/5\%\,\text{CO}_2$. The transfection mixture was then be removed by aspiration, followed by the addition of 25ml of DMEM/10% Fetal Bovine Serum. Cells will then be incubated at $37^{\circ}\text{C}/5\%\,\text{CO}_2$. After 72hr incubation, cells can then be harvested and utilized for analysis.

5 Example 3

TISSUE DISTRIBUTION OF THE DISCLOSED HUMAN GPCRS

Several approaches can be used for determination of the tissue distribution of the GPCRs disclosed herein.

1. Dot-Blot Analysis

Using a commercially available human-tissue dot-blot format, endogenous orphan GPCRs were probed for a determination of the areas where such receptors are localized. cDNA fragments from the GPCRs of Example 1 (radiolabelled) were (or can be) used as the probe: radiolabeled probe was (or can be) generated using the complete receptor cDNA (excised from the vector) using a Prime-It IITM Random Primer Labeling Kit (Stratagene, 15 #300385), according to manufacturer's instructions. A human RNA Master BlotTM (Clontech, #7770-1) was hybridized with the endogenous human GPCR radiolabeled probe and washed under stringent conditions according manufacturer's instructions. The blot was exposed to Kodak BioMaxTM Autoradiography film overnight at -80°C. Results are summarized for several receptors in Table B and C (see Figures 1A and 1B for a grid 20 identifying the various tissues and their locations, respectively). Exemplary dot-blots are provided in Figure 2A and 2B for results derived using hCHN3 and hCHN8, respectively.

TABLE B

ORPHAN GPCR

Tissue Distribution (highest levels, relative to other tissues in the dot-blot)

	hGPCR27	Fetal brain, Putamen, Pituitary gland, Caudate nucleus
	hARE-1	Spleen, Peripheral leukocytes, Fetal spleen
	hPPR1	Pituitary gland, Heart, salivary gland, Small intestine, Testis
	hRUP3	Pancreas
5	hCHN3	Fetal brain, Putamen, Occipital cortex
	hCHN9	Pancreas, Small intestine, Liver
	hCHN10	Kidney, Thryoid

TABLE C

•	ORPHAN GPCR	Tissue Distribution (highest levels, relative to other tissues in the dot-blot)
10	hARE-3	Cerebellum left, Cerebellum right, Testis, Accumbens
	hGPCR3	Corpus collusum, Caudate nucleus, Liver, Heart, Inter- Ventricular Septum
-	hARE-2	Cerebellum left, Cerebellum right, Substantia
	hCHN8	Cerebellum left, Cerebellum right, Kidney, Lung

2. RT-PCR

15 **a. hRUP3**

To ascertain the tissue distribution of hRUP3 mRNA, RT-PCR was performed using hRUP3-specific primers and human multiple tissue cDNA panels (MTC, Clontech) as templates. Taq DNA polymerase (Stratagene) was utilized for the PCR reaction, using the following reaction cycles in a 40ul reaction: 94°C for 2 min; 94°C for 15 sec; 55°C for 30 20 sec; 72°C for 1 min; 72°C, for 10 min. Primers were as follows:

- 5'-GACAGGTACCTTGCCATCAAG-3' (SEQ.ID.NO.: 61; sense)
- 5'-CTGCACAATGCCAGTGATAAGG-3' (SEQ.ID.NO.: 62; antisense).
- 20ul of the reaction was loaded onto a 1% agarose gel; results are set forth in Figure 3.

As is supported by the data of Figure 3, of the 16 human tissues in the cDNA panel utilized (brain, colon, heart, kidney, lung, ovary, pancreas, placenta, prostate, skeleton, small intestine, spleen, testis, thymus leukocyte, and liver) a single hRUP3 band is evident only from the pancreas. Additional comparative analysis of the protein sequence of hRUP3 with 5 other GPCRs suggest that hRUP3 is related to GPCRs having small molecule endogenous ligand such that it is predicted that the endogenous ligand for hRUP3 is a small molecule.

b. hRUP4

RT-PCR was performed using hRUP4 oligo's 8 and 4 as primers and the human multiple tissue cDNA panels (MTC, Clontech) as templates. Taq DNA polymerase 10 (Stratagene) was used for the amplification in a 40ul reaction by the following cycles: 94°C for 30 seconds, 94°C for 10 seconds, 55°C for 30 seconds, 72°C for 2 minutes, and 72°C for 5 minutes with cycles 2 through 4 repeated 30 times.

20 µl of the reaction were loaded on a 1% agarose gel to analyze the RT-PCR products, and hRUP4 mRNA was found expressed in many human tissues, with the strongest 15 expression in heart and kidney. (see, Figure 4). To confirm the authenticity of the PCR fragments, a 300 bp fragment derived from the 5' end of hRUP4 was used as a probe for the Southern Blot analysis. The probe was labeled with ³²P-dCTP using the Prime-It II™ Random Primer Labeling Kit (Stratagene) and purified using the ProbeQuant™ G-50 micro columns (Amersham). Hybridization was done overnight at 42° C following a 12 hr pre-20 hybridization. The blot was finally washed at 65°C with 0.1 x SSC. The Southern blot did confirm the PCR fragments as hRUP4.

c. hRUP5

RT-PCR was performed using the following hRUP5 specific primers:

- 5'-CTGACTTCTTGTTCCTGGCAGCAGCGG-3' (SEQ.ID.NO.: 63; sense)
- 5'-AGACCAGCCAGGCACGCTGAAGAGTG-3' (SEQ.ID.NO.: 64; antisense)

and the human multiple tissue cDNA panels (MTC, Clontech) as templates. Taq DNA 5 polymerase (Stratagene) was used for the amplification in a 40ul reaction by the following cycles: 94°C for 30 sec, 94°C for 10 sec, 62°C for 1.5 min, 72°C for 5 min, and with cycles 2 through 3 repeated 30 times. 20 µl of the reaction were loaded on a 1.5% agarose gel to analyze the RT-PCR products, and hRUP5 mRNA was found expressed only in the peripheral blood leukocytes (*data not shown*).

10 d. hRUP6

RT-PCR was applied to confirm the expression and to determine the tissue distribution of hRUP6. Oligonucleotides used, based on an alignment of AC005871 and GPR66 segments, had the following sequences:

5'-CCAACACCAGCATCCATGGCATCAAG-3' (SEQ.ID.NO.: 73; sense),

15 5'-GGAGAGTCAGCTCTGAAAGAATTCAGG-3' (SEQ.ID.NO.: 74; antisense)
and the human multiple tissue cDNA panels (MTC, Clontech) were used as templates.

PCR was performed using TaqPlus Precision™ polymerase (Stratagene; manufacturing instructions will be followed) in a 40ul reaction by the following cycles: 94°C for 30 sec; 94°C 5 sec; 66°C for 40 sec, 72°C for 2.5 min, and 72°C for 7 min. Cycles 2 through 4 20 were repeated 30 times.

20 ul of the reaction were loaded on a 1.2% agarose gel to analyze the RT-PCR products, and a specific 760bp DNA fragment representing hRUP6 was expressed predominantly in the thymus and with less expression in the heart, kidney, lung, prostate small intestine and testis. (see, Figure 5).

It is intended that each of the patents, applications, and printed publications mentioned in this patent document be hereby incorporated by reference in their entirety.

As those skilled in the art will appreciate, numerous changes and modifications may be made to the preferred embodiments of the invention without departing from the 5 spirit of the invention. It is intended that all such variations fall within the scope of the invention and the claims that follow.

Although a variety of Vectors are available to those in the art, for purposes of utilization for both endogenous and non-endogenous human GPCRs, it is most preferred that the Vector utilized be pCMV. This vector was deposited with the American Type 10 Culture Collection (ATCC) on October 13, 1998 (10801 University Blvd., Manassas, VA 20110-2209 USA) under the provisions of the Budapest Treaty for the International Recognition of the Deposit of Microorganisms for the Purpose of Patent Procedure. The DNA was tested by the ATCC and determined to be. The ATCC has assigned the following deposit number to pCMV: ATCC #203351.

CLAIMS

What is claimed is:

- 1. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 1.
- 2. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 1 comprising SEQ.ID.NO.: 2.
 - 3. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:1.
 - 4. A Host Cell comprising the Plasmid of claim 3.
- 5. A cDNA encoding a human G protein-coupled receptor comprising 10 SEQ.ID.NO.: 3.
 - 6. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 3 comprising SEQ.ID.NO.: 4.
 - 7. A Plasmid comprising a Vector and the cDNA of SEO.ID.NO.:3.
 - 8. A Host Cell comprising the Plasmid of claim 7.
- 9. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 5.
 - 10. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 5 comprising SEQ.ID.NO.: 6.
 - 11. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:5.
- 20 12. A Host Cell comprising the Plasmid of claim 11.
 - 13. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 7.

- 14. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 7 comprising SEQ.ID.NO.: 8.
 - 15. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:7.
 - 16. A Host Cell comprising the Plasmid of claim 15.
- 17. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 9.
- 18. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 9 comprising SEQ.ID.NO.: 10.
 - 19. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:9.
- 10 20. A Host Cell comprising the Plasmid of claim 19.
 - 21. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 11.
 - 22. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 11 comprising SEQ.ID.NO.:12.
- 15 23. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:11.
 - 24. A Host Cell comprising the Plasmid of claim 23.
- 25. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 13.
- 26. A human G protein-coupled receptor encoded by the cDNA of 20 SEQ.ID.NO.: 13 comprising SEQ.ID.NO.: 14.
 - 27. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:13.
 - 28. A Host Cell comprising the Plasmid of claim 27.
 - 29. A cDNA encoding a human G protein-coupled receptor comprising

SEQ.ID.NO.: 15.

- 30. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 15 comprising SEQ.ID.NO.: 16.
 - 31. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:15.
- 5 32. A Host Cell comprising the Plasmid of claim 31.
 - 33. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 17.
- 34. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 17 comprising SEQ.ID.NO.: 18.
- 10 35. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:17.
 - 36. A Host Cell comprising the Plasmid of claim 35.
 - 37. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 19.
- 38. A human G protein-coupled receptor encoded by the cDNA of 15 SEQ.ID.NO.: 19 comprising SEQ.ID.NO.: 20.
 - 39. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:19.
 - 40. A Host Cell comprising the Plasmid of claim 39.
- 41. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 21.
- 20 42. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 21 comprising SEQ.ID.NO.: 22.
 - 43. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:21.
 - 44. A Host Cell comprising the Plasmid of claim 43.

- 45. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 23.
- 46. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 23 comprising SEQ.ID.NO.: 24.
- 5 47. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.: 23.
 - 48. A Host Cell comprising the Plasmid of claim 47.
- 49. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 25.
- 50. A human G protein-coupled receptor encoded by the cDNA of 10 SEQ.ID.NO.: 25 comprising SEQ.ID.NO.: 26.
 - 51. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:25.
 - 52. A Host Cell comprising the Plasmid of claim 51.
- 53. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 27.
- 54. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 27 comprising SEQ.ID.NO.: 28.
 - 55. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:27.
 - 56. A Host Cell comprising the Plasmid of claim 55.
- 57. A cDNA encoding a human G protein-coupled receptor comprising 20 SEQ.ID.NO.: 29.
 - 58. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 29 comprising SEQ.ID.NO.: 30.
 - 59. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:29.

- 60. A Host Cell comprising the Plasmid of claim 59.
- 61. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 31.
- 62. A human G protein-coupled receptor encoded by the cDNA of 5 SEQ.ID.NO.: 31 comprising SEQ.ID.NO.: 32.
 - 63. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:31.
 - 64. A Host Cell comprising the Plasmid of claim 63.
- 65. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 33.
- 10 66. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 33 comprising SEQ.ID.NO.: 34.
 - 67. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:33.
 - 68. A Host Cell comprising the Plasmid of claim 67.
- 69. A cDNA encoding a human G protein-coupled receptor comprising 15 SEQ.ID.NO.: 35.
 - 70. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 35 comprising SEQ.ID.NO.: 36.
 - 71. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:35.
 - 72. A Host Cell comprising the Plasmid of claim 71.
- 20 73. A cDNA encoding a human G protein-coupled receptor comprising SEQ.ID.NO.: 37.
 - 74. A human G protein-coupled receptor encoded by the cDNA of SEQ.ID.NO.: 37 comprising SEQ.ID.NO.: 38.

- 75. A Plasmid comprising a Vector and the cDNA of SEQ.ID.NO.:37.
- 76. A Host Cell comprising the Plasmid of claim 75.

-	- 11	· ·																	
	∞	Medulla	Ohlongata	Colougada		Stomooh	החוושרוו		Mammary	Closed	Clalid	Bone	N Comment	IVIALIUW					
L	_	Hippocampus Medulla		Spinal Cord		Prostate			Salivary	Gland	Oraliu	Lymph	Node	TAORIC		Fetal	7	Lung	
y	0	Frontal	Cortex	A		Uterus			Thyroid			Peripheral	Lenkocyte	21		Fetal	Thymne	contrigity	
.	, ,	Cerabral	Cortex	Thalamus		Bladder			Adrenal	Gland	I	Thymus				Fetal	Shleen	Tracella	
4	Constant	Cerebellum		Temporal	Cortex	Colon		٠. د	Pituitary		7	Spieen		Placenta	ŗ	retal	Liver		
3	Candata		Nucleus	Substantia	Nigra	Skeletal	Muscle	D	rancreas		Crac 11	Siliali	Intestine	Trachea	17-7-1	retai	Kidney		
2	Amvodala	· ··· · J & daild -		Putamen	-	Aorta		Jr. 1024	Oval y		Tiver	7,17		- Lung	Total.	rcial	Heart		
1				Occipital	Cortex	Heart		Tectic	CINCAT		Kidney	Communication		Appendix	Fefal	ı vial	Brain		
	A		,	<u>n</u>		<u>ပ</u>					ĹΤ			ľ,	٢)		H	

FIG. 1A

SUBSTITUTE SHEET (RULE 26)

	[2]			Ť		\exists			Γ		Τ	·		2 /	<u> </u>		Т				T		
•	Ξ	Fetal	Drain	Teta!	2 7	Heart	Fetal	Kidney	Fetal	Liver	Fetal	Spleen		Fetal	Thymus		Fetal	Lung)				-
-	10	Leukemia	HI 60	Hela S3			Leukemia	K562	Leukemia	MOLT-4	Burkitt's	Lymphoma	Raii	Burkitt's	Lymphona	Daudi	Colorectal	Adenocarcinoma	SW480		Timo	Carcinenta	A549
	6	Liver		Pancreas		A demonstrated	Adrenal	Cland	Thyroid		Salivary	Gland		Mammary	Gland				•				
	∞	Lung	······································	Placenta		Bladder	חשות		Uterus		Prostate			Testis		•	Ovary						
	7	Kidney	,	Skeletal	Muscle	Shleen	i Nich		Ihymus		Peripheral	Leukocyte		Lymph	Node		Bone	Marrow			Trachea		
	9	Colon	Transverse	Colon	Desending	Rechim			_									-					
	\sim	Esophagus		Stornach		Duodenum		1	nejmum	ī	lleum			Ilocecum			Appendix				Colon	Ascending	
,	+	Heart		Aorta		Atrium	I eff	, A	11111111111111111111111111111111111111	יי אניין א	ventrole	Ħ		Ventricle	III Marie		Inter 17	venincular	winds.		Apex of	the Heart	
3	ر ا	Substantia	Nigra	Accumbens		Thalamus		Pitnitary,	Gland	Chinol		ם מ											
2	1 -	Cerebellum	III .		Right	Corpus	Callosum	Amvodala	000	Candata	Niclens	STORY I	1.1	Suduexoddira		Machilla	Ohlonom	Colougala		7	rutamen		
1			\perp		Cortex	Frontal	Cortex	Parietal	Lobe	Occinital	Cortex		Terranour	Cortex		G Paracentral	Gyms of	Cerebral	Cortex	Done	9 D T		
·	△	ζ	P	Δ		<u>၂</u>		Ω		回			Ĺī	٠		C				垣	7]

FIG. 18

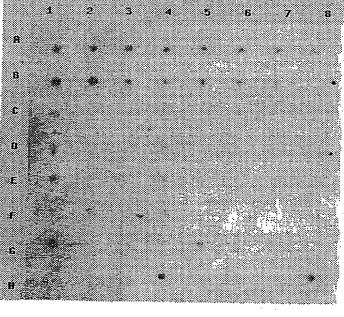


FIG. 2A

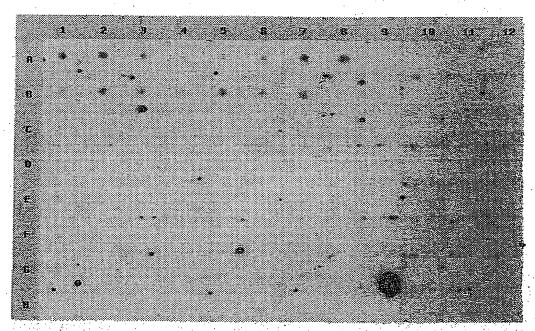


FIG. 2B

SUBSTITUTE SHEET (RULE 26)

H20
Thymus
Testis
Spleen
Small Intestine
Skeletal Muscle

Placenta

Pancreas

Ovary Lung Liver Leukocyte Kidney

Heart Colon Brain

H2O
Thymus
Testis
Spleen
Small intestine
Skeletal Muscle
Prostate
Placenta
1 kbt Lad.
Pancreas
Ovary
Lung
Liver
Leukocyte
Kidney
Heart
Colon
Brain

Brain Colon Heart Kidney Lung Ovary Pancreas Placenta

Skeletal Muscle Small Intestine

Spleen Testis

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FIG. 4

SEQUENCE LISTING

•	_	ד א סיטואיטיט ז	INFORMATION:
١	·	GENERAL	TMLOKMALION.

(i) APPLICANT: Chen, Ruoping
Dang, Huong T.
Liaw, Chen W.
Lin, I-Lin

- (ii) TITLE OF INVENTION: Human Orphan G Protein-Coupled Receptors
- (iii) NUMBER OF SEQUENCES: 74
- 10 (iv) CORRESPONDENCE ADDRESS:
 - (A) ADDRESSEE: Arena Pharmaceuticals, Inc.
 - (B) STREET: 6166 Nancy Ridge Drive
 - (C) CITY: San Diego
 - (D) STATE: CA
- 15 (E) COUNTRY: USA
 - (F) ZIP: 92121
 - (v) COMPUTER READABLE FORM:
 - (A) MEDIUM TYPE: Floppy disk
 - (B) COMPUTER: IBM PC compatible
- 20 (C) OPERATING SYSTEM: PC-DOS/MS-DOS
 - (D) SOFTWARE: PatentIn Release #1.0, Version #1.30
 - (vi) CURRENT APPLICATION DATA:
 - (A) APPLICATION NUMBER: US
 - (B) FILING DATE:
- 25 (C) CLASSIFICATION:

- (viii) ATTORNEY/AGENT INFORMATION:
 - (A) NAME: Burgoon, Richard P.
 - (B) REGISTRATION NUMBER: 34,787
- (ix) TELECOMMUNICATION INFORMATION:
 - (A) TELEPHONE: (858)453-7200
 - (B) TELEFAX: (858)453-7210
- (2) INFORMATION FOR SEQ ID NO:1:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1260 base pairs
- 35 (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA (genomic)
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:
- 40 ATGGTCTTCT CGGCAGTGTT GACTGCGTTC CATACCGGGA CATCCAACAC AACATTTGTC 60

GTGTATGAAA ACACCTACAT GAATATTACA CTCCCTCCAC CATTCCAGCA TCCTGACCTC 120 AGTCCATTGC TTAGATATAG TTTTGAAACC ATGGCTCCCA CTGGTTTGAG TTCCTTGACC 180 GTGAATAGTA CAGCTGTGCC CACAACACCA GCAGCATTTA AGAGCCTAAA CTTGCCTCTT 240 CAGATCACCC TTTCTGCTAT AATGATATTC ATTCTGTTTG TGTCTTTTCT TGGGAACTTG 300 5 GTTGTTTGCC TCATGGTTTA CCAAAAAGCT GCCATGAGGT CTGCAATTAA CATCCTCCTT 360 GCCAGCCTAG CTTTTGCAGA CATGTTGCTT GCAGTGCTGA ACATGCCCTT TGCCCTGGTA 420 ACTATTCTTA CTACCCGATG GATTTTTGGG AAATTCTTCT GTAGGGTATC TGCTATGTTT 480 TTCTGGTTAT TTGTGATAGA AGGAGTAGCC ATCCTGCTCA TCATTAGCAT AGATAGGTTC 540 CTTATTATAG TCCAGAGGCA GGATAAGCTA AACCCATATA GAGCTAAGGT TCTGATTGCA 600 10 GTTTCTTGGG CAACTTCCTT TTGTGTAGCT TTTCCTTTAG CCGTAGGAAA CCCCGACCTG 660 CAGATACCTT CCCGAGCTCC CCAGTGTGTG TTTGGGTACA CAACCAATCC AGGCTACCAG 720 GCTTATGTGA TTTTGATTTC TCTCATTTCT TTCTTCATAC CCTTCCTGGT AATACTGTAC 780 TCATTTATGG GCATACTCAA CACCCTTCGG CACAATGCCT TGAGGATCCA TAGCTACCCT 840 GAAGGTATAT GCCTCAGCCA GGCCAGCAAA CTGGGTCTCA TGAGTCTGCA GAGACCTTTC 900 15 CAGATGAGCA TTGACATGGG CTTTAAAACA CGTGCCTTCA CCACTATTTT GATTCTCTTT 960 GCTGTCTTCA TTGTCTGCTG GGCCCCATTC ACCACTTACA GCCTTGTGGC AACATTCAGT1020 AAGCACTTTT ACTATCAGCA CAACTTTTTT GAGATTAGCA CCTGGCTACT GTGGCTCTGC1080 TACCTCAAGT CTGCATTGAA TCCGCTGATC TACTACTGGA GGATTAAGAA ATTCCATGAT1140 20 GCTTGCCTGG ACATGATGCC TAAGTCCTTC AAGTTTTTGC CGCAGCTCCC TGGTCACACA1200 AAGCGACGGA TACGTCCTAG TGCTGTCTAT GTGTGTGGGG AACATCGGAC GGTGGTGTGA1260

(3) INFORMATION FOR SEQ ID NO:2:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 419 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:
- Met Val Phe Ser Ala Val Leu Thr Ala Phe His Thr Gly Thr Ser Asn

	Thi	r Thi	r Phe	e Va 20	l Va	1 : T y	r Gli	u Ası	n Th: 25	т Ту	r Me	t Ası	n Il	e Th:	r Le	ı Pr
	Pro	Pro	35	e Gli	n Hi	s Pro	a Ası	Let 40	ı Sei	r Pro	· Det	ı Leı	ı Arç 45	д Ту	r Sei	r Ph
5	Glu	Thr 50	Met	: Ala	a Pro	o Thi	55	, Leι	ı Sei	Sez	r Let	Thi 60	· Val	l Ası	ı Sei	Th
	Ala 65	∵Va1	Pro	Thi	Th:	70	Ala	a Ala	≀ Ph∈	Lys	S Ser	Lei	ı Asr	ı Lei	ı Pro	Let 80
10	Gln	Ile	Thr	Leu	Ser 85	Ala	ılle	Met	Ile	Phe 90	Ile	Leu	Phe	· Val	Ser 95	Phe
	Leu	Gly	Asn	Leu 100	Val	. Val	Cys	Leu	Met 105		Tyr	Gln	Lys	Ala 110		Met
	Arg	Ser	Ala 115	Ile	Asn	Ile	Leu	Leu 120	Ala	Ser	Leu	Ala	Phe 125		Asp	Met
15	Leu	Leu 130	Ala	Val	Leu	Asn	Met 135	Pro	Phe	Ala	Leu	Val 140		Ile	Leu	Thr
	Thr 145	Arg	Trp	Ile	Phe	Gly 150	Lys	Phe	Phe	Суз	Arg 155	Val	Ser	Ala	Met	Phe 160
20					165		Glu			170		•			175	
				180			Ile		185		•			190		. ,
			195				Ile	200:					205			
25		210				4.7	Val 215				•	220				
	225					230	Phe	•			235					240
30	Ala				245		* ,			250					255	
	Val			260	•	. *			265					270		
0.7	Ala		275					280		•			285			
35		290					295					300				
•	Asp 1	Met (Gly 1	Phe :	Lys '	Thr .	Arg i	Ala :	Phe '	Thr '	Thr	Ile :	Leu	Ile	Leu :	Phe

	305					310					315					320
	Ala	Val	Phe	Ile	Val 325	Cys	Trp	Ala		Phe 330		Thr	Tyr	Ser	Leu 335	Val
5	Ala	Thr	Phe	Ser 340	Lys	His	Phe	Tyr	Tyr 345	Gln	His	Asn	Phe	Phe 350		Ile
	Ser	Thr	Trp 355	Leu	Leu	Trp	Leu	Cys 360	Tyr	Leu	Lys		Ala 365	Leu	Asn	Pro
	Leu	Ile 370	Tyr	Tyr	Trp	Arg	Ile 375	Lys	Lys	Phe	His	Asp 380		Cys	Leu	Asp
10	Met. 385	Met	Pro	Lys	Ser	Phe 390	Lys	Phe	Leu	Pro	Gln 395		Pro	Gly		Thr 400
	Lys	Arg	Arg	Ile	Arg 405	Pro	Ser	Ala	Val	Tyr 410	Val	Cys	Gly	Glu	His 415	Arg
15	Thr	Val	Val				. •									•
(4)	INFOR	RMAT'I	ON F	OR S	EO T	אר ד										

- - (i) SÉQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1119 base pairs
 - (B) TYPE: nucleic acid
- 20 (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA (genomic)
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:
- ATGTTAGCCA ACAGCTCCTC AACCAACAGT TCTGTTCTCC GGTGTCCTGA CTACCGACCT 60 25 ACCCACCGCC TGCACTTGGT GGTCTACAGC TTGGTGCTGG CTGCCGGGCT CCCCCTCAAC 120 GCGCTAGCCC TCTGGGTCTT CCTGCGCGCG CTGCGCGTGC ACTCGGTGGT GAGCGTGTAC 180 ATGTGTAACC TGGCGGCCAG CGACCTGCTC TTCACCCTCT CGCTGCCCGT TCGTCTCTCC 240 TACTACGCAC TGCACCACTG GCCCTTCCCC GACCTCCTGT GCCAGACGAC GGGCGCCATC 300 TTCCAGATGA ACATGTACGG CAGCTGCATC TTCCTGATGC TCATCAACGT GGACCGCTAC 360 30 GCCGCCATCG TGCACCCGCT GCGACTGCGC CACCTGCGGC GGCCCCGCGT GGCGCGGCTG 420 CTCTGCCTGG GCGTGTGGGC GCTCATCCTG GTGTTTGCCG TGCCCGCCGC CCGCGTGCAC 480 AGGCCCTCGC GTTGCCGCTA CCGGGACCTC GAGGTGCGCC TATGCTTCGA GAGCTTCAGC 540 GACGAGCTGT GGAAAGGCAG GCTGCTGCCC CTCGTGCTGC TGGCCGAGGC GCTGGGCTTC 600

CTGCTGCCCC TGGCGGCGT GGTCTACTCG TCGGGCCGAG TCTTCTGGAC GCTGGCGCGC 660

CCCGACGCCA CGCAGAGCCA GCGGCGGCGG AAGACCGTGC GCCTCCTGCT GGCTAACCTC 720

GTCATCTTCC TGCTGTGCTT CGTGCCCTAC AACAGCACGC TGGCGGTCTA CGGGCTGCTG 780

CGGAGCAAGC TGGTGGCGG CAGCGTGCCT GCCCGCGATC GCGTGCGCG GGTGCTGATG 840

5 GTGATGGTGC TGCTGGCCGG CGCCAACTGC GTGCTGGACC CGCTGGTGA CTACTTTAGC 900

GCCGACGAGC TCCGCAACAC CCTGCGCGGC CTGGGCACTC CGCACCGGGC CAGGACCTCG 960

GCCACCAACG GGACGCGGC CAGTCAGGG CTGCTCCGAAA GGTCCGCCGT CACCACCGAC1020

GCCACCAGGC CGGATGCCGC CAGTCAGGGG CTGCTCCGAC CCTCCGACTC CCACTCTCTG1080

TCTTCCTTCA CACAGTGTCC CCAGGATTCC GCCCTCTGA

10 (5) INFORMATION FOR SEQ ID NO:4:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 372 amino acids
 - (B) TYPE: amino acid
- (C) STRANDEDNESS:
- 15 (D) TOPOLOGY: not relevant
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:
 - Met Leu Ala Asn Ser Ser Ser Thr Asn Ser Ser Val Leu Pro Cys Pro 1 5 10 15
- 20 Asp Tyr Arg Pro Thr His Arg Leu His Leu Val Val Tyr Ser Leu Val 20 25 30
 - Leu Ala Ala Gly Leu Pro Leu Asn Ala Leu Ala Leu Trp Val Phe Leu 35 40 45
- Arg Ala Leu Arg Val His Ser Val Val Ser Val Tyr Met Cys Asn Leu 25 50 55 60
 - Ala Ala Ser Asp Leu Leu Phe Thr Leu Ser Leu Pro Val Arg Leu Ser 65 70 75 80
 - Tyr Tyr Ala Leu His His Trp Pro Phe Pro Asp Leu Leu Cys Gln Thr 85 90 95
- Thr Gly Ala Ile Phe Gln Met Asn Met Tyr Gly Ser Cys Ile Phe Leu 100 105 110
 - Met Leu Ile Asn Val Asp Arg Tyr Ala Ala Ile Val His Pro Leu Arg 115 120 125

	Le	u Ar 13	g Hi O	s Le	u Arg	J Arg	135	Arg	y Val	Ala	Arg	140		ı Cys	Leu	ı Gl
	Va. 14	1 Tr;	p Al	a Lei	u Ile	150	ı Val	Phe	e Ala	·Val	Pro 155		Ala	a Arg	Val	. Hi.
5	Ar	g Pro	o Se	r Arg	7 Cys 165	Arg	тух	Arg	Asr	Leu 170		Val	Arg	Leu	°Cys	
	Glı	ı Ser	r Phe	2 Sei	Asp	Glu	Leu	Trp	Lys 185	Gly	Arg	Leu	Leu	Pro 190		Va.
10	Le	ı Leı	1 Ala 199	a Glu	ı Ala	Leu	Gly	Phe 200		Leu	Pro	Leu	Ala 205		Val	Val
	Туг	210	Ser	Gly	Arg	Val	Phe 215	Trp	Thr	Leu	Ala	Arg 220	Pro	Asp	Ala	Thr
	Glr 225	ser	Glr	Arg	Arg	Arg 230	Lys	Thr	Val	Arg	Leu 235	Leu	Leu	Àla	Asn	Leu 240
15	Val	. Ile	Phe	. Leu	Leu 245	Cys	Phe	Val	Pro	Tyr 250	Asn	Ser	Thr	Leu	Ala 255	Val
	Tyr	. GlA	Leu	Leu 260	Arg	Ser	Lys	Leu	Val 265	Ala	Ala	Ser	Val	Pro 270	Ala	Arg
20	Asp	Arg	Val 275	Arg	Gly	Val	Leu	Met 280	Val	Met	Val	Leu	Leu 285	Ala	Gly	Ala
	Asn	Cys 290	Val	Leu	Asp	Pro	Leu 295	Val	Tyr	Tyr		Ser 300	Ala	Glu	Gly	Phe
	Arg 305	Asn	Thr	Leu	Arg	Gly 310	Leu	Gly	Thr	Pro	His 315	Arg	Ala	Arg	Thr	Ser 320
25	Ala	Thr	Asn	Gly	Thr 325	Arg	Ala	Ala	Leu	Ala 330	Gln	Ser	Glu	Arg	Ser 335	Ala
	Val	Thr	Thr	Asp 340	Ala	Thr		Pro		Ala	Ala	Ser	Gln	Gly 350	Leu	Leu
0	Arg	Pro	Ser 355	Asp	Ser	His		Leu 360	Ser	Ser :	Phe '		Gln 365	Сув	Pro	Gln
	Asp	Ser 370	Ala	Leu				-								:

(6) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1107 base pairs (B) TYPE: nucleic acid

 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:5: ATGGCCAACT CCACAGGGCT GAACGCCTCA GAAGTCGCAG GCTCGTTGGG GTTGATCCTG GCAGCTGTCG TGGAGGTGGG GGCACTGCTG GGCAACGGCG CGCTGCTGGT CGTGGTGCTG 120 5 CGCACGCCGG GACTGCGCGA CGCGCTCTAC CTGGCGCACC TGTGCGTCGT GGACCTGCTG 180 GCGGCCGCCT CCATCATGCC GCTGGGCCTG CTGGCCGCAC CGCCGCCCGG GCTGGGCCGC 240 GTGCGCCTGG GCCCCGCGC ATGCCGCGC GCTCGCTTCC TCTCCGCCGC TCTGCTGCCG 300 GCCTGCACGC TCGGGGTGGC CGCACTTGGC CTGGCACGCT ACCGCCTCAT CGTGCACCCG 360 CTGCGGCCAG GCTCGCGGCC GCCGCCTGTG CTCGTGCTCA CCGCCGTGTG GGCCGCGGCG 420 10 GGACTGCTGG GCGCGCTCTC CCTGCTCGGC CCGCCGCCCG CACCGCCCCC TGCTCCTGCT 480 CGCTGCTCGG TCCTGGCTGG GGGCCTCGGG CCCTTCCGGC CGCTCTGGGC CCTGCTGGCC 540 TTCGCGCTGC CCGCCCTCCT GCTGCTCGGC GCCTACGGCG GCATCTTCGT GGTGGCGCGT 600 CGCGCTGCCC TGAGGCCCCC ACGGCCGGCG CGCGGGTCCC GACTCCGCTC GGACTCTCTG 660 GATAGCCGCC TTTCCATCTT GCCGCCGCTC CGGCCTCGCC TGCCCGGGGG CAAGGCGGCC 720 15 CTGGCCCCAG CGCTGGCCGT GGGCCAATTT GCAGCCTGCT GGCTGCCTTA TGGCTGCGCG 780 TGCCTGGCGC CCGCAGCGCG GGCCGCGGAA GCCGAAGCGG CTGTCACCTG GGTCGCCTAC 840 TCGGCCTTCG CGGCTCACCC CTTCCTGTAC GGGCTGCTGC AGCGCCCCGT GCGCTTGGCA 900 CTGGGCCGCC TCTCTCGCCG TGCACTGCCT GGACCTGTGC GGGCCTGCAC TCCGCAAGCC 960 TGGCACCCGC GGGCACTCTT GCAATGCCTC CAGAGACCCC CAGAGGGCCC TGCCGTAGGC1020 20 CCTTCTGAGG CTCCAGAACA GACCCCCGAG TTGGCAGGAG GGCGGAGCCC CGCATACCAG1080 GGGCCACCTG AGAGTTCTCT CTCCTGA 1107
 - (7) INFORMATION FOR SEO ID NO:6:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 368 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS:
 - (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: protein
- (xi) SEQUENCE DESCRIPTION: SEO ID NO:6:

	Me	t Al	a As	n Se	r Thi	r Gly	y Let	ı Asr	a Ala	Ser 10	Glı	ı Val	L Ala	a Gly	7 Set	r Le
·	Gly	y Le	u Il	e Le: 20	ı Ala	a Ala	a Val	. Val	Glu 25	ı Val	Gl3	/ Ala	Let	ı Let 30	ı Gly	/ Ası
5	Gl	/ Ala	a Let	u Let	ı Val	. Val	. Val	Leu 40	Arg	Thr	Pro	Gly	Leu 45	ı Arç	as.	Ala
	Leu	ту 50	r Lei	ı Ala	a His	Lev	Cys 55	Val	Val	Asp	Leu	Leu 60	Ala	Ala	Àla	Se:
10	Il∈ 65	. Met	Pro) Lev	Gly	Leú 70	i Leu	Ala	Ala	Pro	Pro 75	Pro	Gly	Leu	. Gly	Arg
	Val	. Arg	J Leu	ı Gly	Pro 85	Ala	Pro	Cys	Arg	Ala 90	Ala	Arg	Phe	Leu	Ser 95	Ala
	Ala	Leu	ı Lev	100	Ala	Cys	Thr	Leu	Gly 105		Ala	Ala	Leu	Gly	Leu	Ala
15	Arg	Туг	115	Leu	Ile	Val	His	Pro 120	Leu	Arg	Pro	Gly	Ser 125		Pro	Pro
·	Pro	Val 130	Leu	Val	Leu	Thr	Ala 135	Val	Trp	Ala	Ala	Ala 140	Gly	Leu	Leu	Gly
20	Ala 145	Leu	Ser	Leu	Leu	Gly 150	Pro	Pro	Pro	Ala	Pro 155	Pro	Pro	Ala	Pro	Ala 160
	Arg	Cys	Ser	Val	Leu 165	Ala	Gly	Gly	Leu	Gly 170		Phe	Arg	Pro	Leu 175	Trp
	Ala	Leu	Leu	Ala 180	Phe	Ala	Leu	Pro	Ala 185	Leu	Leu	Leu	Leu	Gly 190	Ala	Tyr
25	Gly	Gly	Ile 195	Phe	Val	Val	Ala	Arg 200	Arg	Ala	Ala	Leu	Arg 205	Pro	Pro	Arg
	Pro	Ala 210	Arg	Gly	Ser	Arg	Leu 215	Arg	Ser	Asp	Ser	Leu 220	Asp	Ser	Arg	Leu
30	Ser 225	Ile	Leu	Pro	Pro	220	Arg •	Pro	Arg	Leu	Pro 235		Gly	Lys	Ala	Ala 240
	Leu	Ala	Pro	Ala	Leu 245	Ala	Val	Gly		Phe : 250	Ala	Ala	Сув	Trp	Leu 255	Pro
	Tyr	Gly	Cys	Ala 260	Cys	Leu	Ala :		Ala . 265		Arg	Ala .		Glu 270	Ala	Glu
35	Ala	Ala	Val 275	Thr	Trp	Val	Ala	Tyr : 280	Ser 2	Ala	Phe		Ala 285	His	Pro	Phe
	Leu	Tyr	Gly	Leu	Leu (Gln .	Arg 1	Pro 1	Val 1	Arg	Leu i	Ala :	Leu	Gly .	Arg	Leu

290 295 300

Ser Arg Arg Ala Leu Pro Gly Pro Val Arg Ala Cys Thr Pro Gln Ala 305 310 315 320

Trp His Pro Arg Ala Leu Leu Gln Cys Leu Gln Arg Pro Pro Glu Gly
325 330 • 335

Pro Ala Val Gly Pro Ser Glu Ala Pro Glu Gln Thr Pro Glu Leu Ala 340 345 350

Gly Gly Arg Ser Pro Ala Tyr Gln Gly Pro Pro Glu Ser Ser Leu Ser 355 360 365

10 (8) INFORMATION FOR SEQ ID NO:7:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1008 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
- 15 (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA (genomic)
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

ATGGAATCAT CATTCTCATT TGGAGTGATC CTTGCTGTCC TGGCCTCCT CATCATTGCT 60
ACTAACACCA TAGTGGCTGT GGCTGTGCTG CTGTTGATCC ACAAGAATGA TGGTGTCAGT 120
ACTACTCACCAG ACCAGCTCTC GGCCTCTCT CGGCCCACCAC AGAAGACCCT GTGCAGCCTG 240
ACGGATGGCAT TTGTCACTTC CACCCCAGCT GCCTCTCT CGCCCACCAC AGAAGACCCT GACGTGCACC 300
ACGGATGGCAT ACCTTGCCAT CAAGCCAGCCC TTCCGCTACT TGAAGACCAT GACTGGGTCC 300
ACGGCCGGGG CCTGCATTCC CGGCCTGTCC TGCGCTACT ACCTCATTGG CTTCCCCCA 420
ACCTCGGAATCC CCATTGCCAT GCAGACTGC TACCAAAGGGC AGTGCAGCTT CTTCCCCAC ACCTCTCC ACCTCCCA ACCTCTCCCA ACCTCTCCCA ACCTCTCCCA ACCTCTCCCA ACCTCTCCCA ACCTCTCCA ACCTCTCCCA ACCTCTCCCA ACCTCTCCCA ACCTCTCCCA ACCTCTCCCA ACCTCTCCA ACCTCTCCCA ACCTCTCCA ACCTCTCA ACCTCTCA

GAACGGTACC TGTGGCTGCT CGGCGTGGGC AACTCCCTGC TCAACCCACT CATCTATGCC 840

TATTGGCAGA AGGAGGTGCG ACTGCAGCTC TACCACATGG CCCTAGGAGT GAAGAAGGTG 900 CTCACCTCAT TCCTCCTCTT TCTCTCGGCC AGGAATTGTG GCCCAGAGAG GCCCAGGGAA 960 AGTTCCTGTC ACATCGTCAC TATCTCCAGC TCAGAGTTTG ATGGCTAA 1008

- (9) INFORMATION FOR SEQ ID NO:8:
- 5 (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 335 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS:
 - (D) TOPOLOGY: not relevant
- 10 (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:
 - Met Glu Ser Ser Phe Ser Phe Gly Val Ile Leu Ala Val Leu Ala Ser 1 5 10 15
- Leu Ile Ile Ala Thr Asn Thr Leu Val Ala Val Ala Val Leu Leu Leu 15 20 25 30
 - Ile His Lys Asn Asp Gly Val Ser Leu Cys Phe Thr Leu Asn Leu Ala 35 40 45
 - Val Ala Asp Thr Leu Ile Gly Val Ala Ile Ser Gly Leu Leu Thr Asp
 50 55 60
- 20 Gln Leu Ser Ser Pro Ser Arg Pro Thr Gln Lys Thr Leu Cys Ser Leu 65 70 75 80
 - Arg Met Ala Phe Val Thr Ser Ser Ala Ala Ala Ser Val Leu Thr Val
- Met Leu Ile Thr Phe Asp Arg Tyr Leu Ala Ile Lys Gln Pro Phe Arg 25 100 105 110
 - Tyr Leu Lys Ile Met Ser Gly Phe Val Ala Gly Ala Cys Ile Ala Gly 115 120 125
 - Leu Trp Leu Val Ser Tyr Leu Ile Gly Phe Leu Pro Leu Gly Ile Pro
 130 140
- Met Phe Gln Gln Thr Ala Tyr Lys Gly Gln Cys Ser Phe Phe Ala Val
 - Phe His Pro His Phe Val Leu Thr Leu Ser Cys Val Gly Phe Phe Pro 165 170 175
 - Ala Met Leu Leu Phe Val Phe Phe Tyr Cys Asp Met Leu Lys Ile Ala 180 185 190

- Gln Leu Tyr His Met Ala Leu Gly Val Lys Lys Val Leu Thr Ser Phe 290 295 300
- Leu Leu Phe Leu Ser Ala Arg Asn Cys Gly Pro Glu Arg Pro Arg Glu 305 310 315 320
 - Ser Ser Cys His Ile Val Thr Ile Ser Ser Ser Glu Phe Asp Gly 325 330 335
 - (10) INFORMATION FOR SEQ ID NO:9:
- 20 (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1413 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- 25 (ii) MOLECULE TYPE: DNA (genomic)
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:
- ATGGACACTA CCATGGAAGC TGACCTGGGT GCCACTGGCC ACAGGCCCC CACAGAGCTT 60
 GATGATGAGG ACTCCTACCC CCAAGGTGGC TGGGACACGG TCTTCCTGGT GGCCCTGCTG 120
 CTCCTTGGGC TGCCAGCCAA TGGGTTGATG GCGTGGCTG CCGGCTCCCA GGCCCGCCAT 180
 30 GGAGCTGGCA CGCGTTCCA GATCCTAGAG ATCCGGCATG CCCTCTCTGA CTTCTTGTTC 240
 ACAGCTGCCT GCCGCTTCCA GATCCTAGAG ATCCGGCATG GGGGACACTG GCCGCTGGGG 300
 CTGCTGGCCG CCCTCAGCCT CGACCCTCC CGGCCTCTC CGGCCTCTC CGGCCTCTC ACAGCCTGC CTGCTGGCC CCGCCCCACA CTGGTACCCT 420
 GGGCACCGCC CAGTCCGCC GCCCCTCTGG GTCTCGCCC GTGTCCGCC GCTGGCCACA 480

CTCTTCAGCG TGCCCTGGCT GGTCTTCCCC GAGGCTGCCG TCTGGTGGTA CGACCTGGTC 540 ATCTGCCTGG ACTTCTGGGA CAGCGAGGAG CTGTCGCTGA GGATGCTGGA GGTCCTGGGG 600 GGCTTCCTGC CTTTCCTCCT GCTGCTCGTC TGCCACGTGC TCACCCAGGC CACAGCCTGT 660 CGCACCTGCC ACCGCCAACA GCAGCCCGCA GCCTGCCGGG GCTTCGCCCG TGTGGCCAGG 720 5 ACCATTCTGT CAGCCTATGT GGTCCTGAGG CTGCCCTACC AGCTGGCCCA GCTGCTCTAC 780 CTGGCCTTCC TGTGGGACGT CTACTCTGGC TACCTGCTCT GGGAGGCCCT GGTCTACTCC 840 GACTACCTGA TCCTACTCAA CAGCTGCCTC AGCCCCTTCC TCTGCCTCAT GGCCAGTGCC 900 GACCTCCGGA CCCTGCTGCG CTCCGTGCTC TCGTCCTTCG CGGCAGCTCT CTGCGAGGAG 960 CGGCCGGGCA GCTTCACGCC CACTGAGCCA CAGACCCAGC TAGATTCTGA GGGTCCAACT1020 10 CTGCCAGAGC CGATGGCAGA GGCCCAGTCA CAGATGGATC CTGTGGCCCA GCCTCAGGTG1080 AACCCCACAC TCCAGCCACG ATCGGATCCC ACAGCTCAGC CACAGCTGAA CCCTACGGCC1140 CAGCCACAGT CGGATCCCAC AGCCCAGCCA CAGCTGAACC TCATGGCCCA GCCACAGTCA1200 GATTCTGTGG CCCAGCCACA GGCAGACACT AACGTCCAGA CCCCTGCACC TGCTGCCAGT1260 TCTGTGCCCA GTCCCTGTGA TGAAGCTTCC CCAACCCCAT CCTCGCATCC TACCCCAGGG1320 15 GCCCTTGAGG ACCCAGCCAC ACCTCCTGCC TCTGAAGGAG AAAGCCCCAG CAGCACCCCG1380 CCAGAGGCGG CCCCGGGCGC AGGCCCCACG TGA 1413

(11) INFORMATION FOR SEQ ID NO:10:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 468 amino acids
- 20 (B) TYPE: amino acid
 - (C) STRANDEDNESS:
 - (D) TOPOLOGY: not relevant
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:
- 25 Met Asp Thr Thr Met Glu Ala Asp Leu Gly Ala Thr Gly His Arg Pro 1 5 10 15
 - Arg Thr Glu Leu Asp Asp Glu Asp Ser Tyr Pro Gln Gly Gly Trp Asp 20 25 30
- Thr Val Phe Leu Val Ala Leu Leu Leu Gly Leu Pro Ala Asn Gly
 30 35 40 45
 - Leu Met Ala Trp Leu Ala Gly Ser Gln Ala Arg His Gly Ala Gly Thr

		50					55					. 60				
•	Arg 65	g Lei	ı Ala	a Leu	ı Let	1 Le: 70	u Lei	ı Se:	r Le	u Ala	a Lei 75	ı Se:	r Ası	Phe	e Let	2 Ph 80
5	Let	ı Ala	Ala	a Ala	Ala 85	Phe	e Glr	ı Ile	e Let	ı Glı 90	ı Ile	e Aro	g His	Gly	7 Gly 95	/ Hi
	Trp	Pro	Let	1 Gly 100		Ala	a Ala	ı Cys	105		э Туз	туз	: Phe	Lei 110		G1
		. ; .	115					120)				125	٠.		
10		130	• .	ı Leu		. 1	. 135					140)			
	145	i		Pro	٠.	150			:		155				•	160
15				Val	165					170				•	175	
				Val 180					185	-		•		190		
			195					200					205		•	
20		210		His			215					220				
	Gln 225	Gln	Pro	Ala	Ala	Суs 230	Arg	Gly	Phe	Ala	Arg 235	Val	Ala	Arg	Thr	Ile 240
25		•		Tyr	245					250					255	
	Leu	Tyr		Ala 260	Phe	Leu	Trp	Asp	Val 265	Tyr	Ser	Gly	Tyr	Leu 270		Trp
	Glu	Ala	Leu 275	Val	Tyr	Ser	Asp	Tyr 280	Leu	Ile	Leu	Leu	Asn 285	Ser	Сув	Leu
30	Ser	Pro . 290	Phe	Leu	Cys	Leu	Met 295	Ala	Ser	Ala	Asp	Leu 300	Arg	Thr	Leu	Leu
	Arg 305	Ser	Val	Leu		Ser 310	Phe	Ala	Ala	Ala	Leu 315	Cys	Glu	Glu	Arg	Pro 320
35	Gly	Ser	Phe	Thr	Pro 325	Thr	Glu	Pro	Gln	Thr 330	Gln	Leu	Asp		Glu 335	Gly
	Pro	Thr	Leu	Pro (Glu	Pro	Met		Glu 345	Ala	Gln	Ser.		Met 350	Asp	Pro

Val	Ala	Gln	Pro.	Gln	Val	Asn	Pro	Thr	Leu	Gln	Pro	Arq	Ser	Asp	Pro
		355					360					365		-	

- Thr Ala Gln Pro Gln Leu Asn Pro Thr Ala Gln Pro Gln Ser Asp Pro 370 375 380
- 5 Thr Ala Gln Pro Gln Leu Asn Leu Met Ala Gln Pro Gln Ser Asp Ser 385 390 395 400
 - Val Ala Gln Pro Gln Ala Asp Thr Asn Val Gln Thr Pro Ala Pro Ala 405 410 415
- Ala Ser Ser Val Pro Ser Pro Cys Asp Glu Ala Ser Pro Thr Pro Ser 10 420 425 430
 - Ser His Pro Thr Pro Gly Ala Leu Glu Asp Pro Ala Thr Pro Pro Ala 435 440 445
 - Ser Glu Gly Glu Ser Pro Ser Ser Thr Pro Pro Glu Ala Ala Pro Gly
 450 455 460
- 15 Ala Gly Pro Thr 465
- (12) INFORMATION FOR SEQ ID NO:11:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1248 base pairs
- 20 (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA (genomic)
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:
- 25 ATGTCAGGGA TGGAAAAACT TCAGAATGCT TCCTGGATCT ACCAGCAGAA ACTAGAAGAT 60

 CCATTCCAGA AACACCTGAA CAGCACCGAG GAGTATCTGG CCTTCCTCTG CGGACCTCGG 120

 CGCAGCCACT TCTTCCTCCC CGTGTCTGTG GTGTATGTGC CAATTTTGT GGTGGGGGTC 180

 ATTGGCAATG TCCTGGTGTG CCTGGTGATT CTGCAGCACC AGGCTATGAA GACGCCCACC 240

 AACTACTACC TCTTCAGCCT GGCGGTCTCT GACCTCCTGG TCCTGCTCCT TGGAATGCCC 300

 30 CTGGAGGTCT ATGAGATGTG GCGCAACTAC CCTTTCTTGT TCGGGCCCGT GGGCTGCTAC 360

 TTCAAGACGG CCCTCTTTGA GACCGTGTGC TTCGCCTCCA TCCTCAGCAT CACCACCGTC 420

 AGCGTGGAGC GCTACGTGG CATCCTACAC CCGTTCCGCG CCAAACTGCA GAGCACCCGG 480

 CGCCGGGCCC TCAGGATCCT CGGCATCGTC TGGGGCTTCT CTCCCTGCCC 540

AACACCAGCA TCCATGGCAT CAAGTTCCAC TACTTCCCCA ATGGGTCCCT GGTCCCAGGT 600
TCGGCCACCT GTACGGTCAT CAAGCCCATG TGGATCTACA ATTTCATCAT CCAGGTCACC 660
TCCTTCCTAT TCTACCTCCT CCCCATGACT GTCATCAGTG TCCTCTACTA CCTCATGGCA 720
CTCAGACTAA AGAAAGACAA ATCTCTTGAG GCAGATGAAG GGAATGCAAA TATTCAAAGA 780
5 CCCTGCAGAA AATCAGTCAA CAAGATGCTG TTTGTCTTGG TCTTAGTGTT TGCTATCTGT 840
TGGGCCCCGT TCCACATTGA CCGACTCTTC TTCAGCTTTG TGGAGGAGTG GAGTGAATCC 900
CTGGCTGCTG TGTTCAACCT CGTCCATGTG GTGTCAGGTG TCCTCTTCTA CCTGAGCTCA 960
GCTGTCAACC CCATTATCTA TAACCTACTG TCTCGCCGCT TCCAGGCAGC ATTCCAGAAT1020
GTGATCTCTT CTTTCCACAA ACAGTGGCAC TCCCAGCATG ACCCACAGTT GCCACCTGCC1080
10 CAGCGGAACA TCTTCCTGAC AGAATGCCAC TTTGTGGAGC TGACCGAAGA TATAGGTCCC1140
CAATTCCCAT GTCAGTCATC CATGCACAAC TCTCACCTCC CAACAGCCCT CTCTAGTGAA1200
CAGATGTCAA GAACAAACTA TCAAAGCTTC CACTTTAACA AAACCTGA 1248

(13) INFORMATION FOR SEQ ID NO:12:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 415 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS:
 - (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: protein
- 20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:
 - Met Ser Gly Met Glu Lys Leu Gln Asn Ala Ser Trp Ile Tyr Gln Gln 1 5 10 10 15
 - Lys Leu Glu Asp Pro Phe Gln Lys His Leu Asn Ser Thr Glu Glu Tyr
 20 25 30
- Leu Ala Phe Leu Cys Gly Pro Arg Arg Ser His Phe Phe Leu Pro Val 35 40 45
 - Ser Val Val Tyr Val Pro Ile Phe Val Val Gly Val Ile Gly Asn Val 50 55 60
- Leu Val Cys Leu Val Ile Leu Gln His Gln Ala Met Lys Thr Pro Thr 30 65 70 75 80
 - Asn Tyr Tyr Leu Phe Ser Leu Ala Val Ser Asp Leu Leu Val Leu Leu 85 90 95

	Le	u Gl	у Ме	t Pro	o Le O	u Gl	u Va	l Ty	r Gl	u Me 5	t Tr	p Ar	g As	n Ty 11		o Ph
	Le	u Ph	e Gl 11	y Pro 5	o Va	1 G1	у Су:	12	r Pho	e Ly	s Th	r Al	a Le 12		e Gl	u Th
5	Va	1 Cy	s Ph 0	e Ala	a Sei	r Il	e Le:	ı Se	r Ile	e Th	r Th	r Va.		r Va	l _• Gl	u Ar
	Ту: 14!	r Va.	l Al	a Ile	e Let	1 His	s Pro	Phe	≥ Arg	g Ala	a Lys 155		u Gli	n Se	r Thi	160
10	Arc	g Arg	g Ala	a Leu	i Arg	g Ile	e Lev	ı Gly	/ Ile	9 Val		Gly	y Phe	e Sei	r Val	
	Phe	e Sei	. Lei	180	Asn	Thx	Ser	Ile	His 185	Gly	/ Ile	Lys	s Phe	His		? Phe
,	Pro) Asr	1 Gly 195	/ Ser	Leu	val	. Pro	Gly 200		Ala	. Thr	Сув	205		. Ile	ь Гув
15	Pro	210	Trp	Ile	Tyr	Asn	Phe 215	Ile	lle	Glr	val	Thr 220		Phe	: Leu	Phe
	Tyr 225	Leu	Leu	Pro	Met	Thr 230	Val	Ile	Ser	Val	Leu 235		Tyr	Leu	Met	Ala 240
20	Leu	Arg	Leu	. Lys	Lys 245	Asp	Lys	Ser	Leu	Glu 250		Asp	Glu	Gly	Asn 255	Ala
	Asn	Ile	Gln	Arg 260	Pro	Cys	Arg	Lys	Ser 265	Val	Asn	Lys	Met	Leu 270	Phe	Val
	Leu	Val	Leu 275	Val	Phe	Ala	Ile	Cys 280	Trp	Ala	Pro	Phe	His 285	Ile	Asp	Arg
25	Leu	Phe 290	Phe	Ser	Phe	Val	.Glu 295	Glu	Trp	Ser	Glu	Ser 300	Leu	Ala	Ala	Val
. •	Phe 305	Asn	Leu	Val	His	Val 310	Val	Ser	Gly	Val	Phe 315	Phe	Tyr	Leu	Ser	Ser 320
0	Ala	Val	Asn	Pro	Ile 325	Ile	Tyr	Asn	Leu	Leu 330	Ser	Arg	Arg	Phe	Gln 335	Ala
				Asn 340 Gln					345					350		
. · 5			355					360		:			365			
		370		Val			375					380				
	385	ser	Ser	Met	His .	Asn 390	Ser	His	Leu	Pro	Thr 395	Ala.	Leu	Ser	Ser	Glu 400

Gln Met Ser Arg Thr Asn Tyr Gln Ser Phe His Phe Asn Lys Thr 405 410 415

(14) INFORMATION FOR SEQ ID NO:13:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1173 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- 10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

ATGCCAGATA CTAATAGCAC AATCAATTTA TCACTAAGCA CTCGTGTTAC TTTAGCATTT TTTATGTCCT TAGTAGCTTT TGCTATAATG CTAGGAAATG CTTTGGTCAT TTTAGCTTTT 120 GTGGTGGACA AAAACCTTAG ACATCGAAGT AGTTATTTT TTCTTAACTT GGCCATCTCT 180 GACTTCTTTG TGGGTGTGAT CTCCATTCCT TTGTACATCC CTCACACGCT GTTCGAATGG 240 15 GATTTTGGAA AGGAAATCTG TGTATTTTGG CTCACTACTG ACTATCTGTT ATGTACAGCA 300 TCTGTATATA ACATTGTCCT CATCAGCTAT GATCGATACC TGTCAGTCTC AAATGCTGTG 360 TCTTATAGAA CTCAACATAC TGGGGTCTTG AAGATTGTTA CTCTGATGGT GGCCGTTTGG 420 GTGCTGGCCT TCTTAGTGAA TGGGCCAATG ATTCTAGTTT CAGAGTCTTG GAAGGATGAA 480 GGTAGTGAAT GTGAACCTGG ATTTTTTCG GAATGGTACA TCCTTGCCAT CACATCATTC 540 20 TTGGAATTCG TGATCCCAGT CATCTTAGTC GCTTATTTCA ACATGAATAT TTATTGGAGC 600 CTGTGGAAGC GTGATCATCT CAGTAGGTGC CAAAGCCATC CTGGACTGAC TGCTGTCTCT 660 TCCAACATCT GTGGACACTC ATTCAGAGGT AGACTATCTT CAAGGAGATC TCTTTCTGCA 720 TCGACAGAAG TTCCTGCATC CTTTCATTCA GAGAGACAGA GGAGAAAGAG TAGTCTCATG 780 TTTTCCTCAA GAACCAAGAT GAATAGCAAT ACAATTGCTT CCAAAATGGG TTCCTTCTCC 840 25 CAATCAGATT CTGTAGCTCT TCACCAAAGG GAACATGTTG AACTGCTTAG AGCCAGGAGA 900 TTAGCCAAGT CACTGGCCAT TCTCTTAGGG GTTTTTGCTG TTTGCTGGGC TCCATATTCT 960 CTGTTCACAA TTGTCCTTTC ATTTTATTCC TCAGCAACAG GTCCTAAATC AGTTTGGTAT1020 AGAATTGCAT TTTGGCTTCA GTGGTTCAAT TCCTTTGTCA ATCCTCTTTT GTATCCATTG1080 TGTCACAAGC GCTTTCAAAA GGCTTTCTTG AAAATATTTT GTATAAAAAA GCAACCTCTAL140 30 CCATCACAAC ACAGTCGGTC AGTATCTTCT TAA 1173

(15)	INFORMATION	FOR	SEQ	ID	NO:14:
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- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 390 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS:
 - (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: protein
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:
- Met Pro Asp Thr Asn Ser Thr Ile Asn Leu Ser Leu Ser Thr Arg Val 10 1 5 10 15
 - Thr Leu Ala Phe Phe Met Ser Leu Val Ala Phe Ala Ile Met Leu Gly 20 25 30
 - Asn Ala Leu Val Ile Leu Ala Phe Val Val Asp Lys Asn Leu Arg His 35 40 45
- 15 Arg Ser Ser Tyr Phe Phe Leu Asn Leu Ala Ile Ser Asp Phe Phe Val 50 55 60
 - Gly Val Ile Ser Ile Pro Leu Tyr Ile Pro His Thr Leu Phe Glu Trp 65 70 75 80
- Asp Phe Gly Lys Glu Ile Cys Val Phe Trp Leu Thr Thr Asp Tyr Leu 20 85 90 95
 - Leu Cys Thr Ala Ser Val Tyr Asn Ile Val Leu Ile Ser Tyr Asp Arg 100 105 110
 - Tyr Leu Ser Val Ser Asn Ala Val Ser Tyr Arg Thr Gln His Thr Gly
 115 120 125
- Val Leu Lys Ile Val Thr Leu Met Val Ala Val Trp Val Leu Ala Phe
 130 135 140
 - Leu Val Asn Gly Pro Met Ile Leu Val Ser Glu Ser Trp Lys Asp Glu
 145 150 155 160
- Gly Ser Glu Cys Glu Pro Gly Phe Phe Ser Glu Trp Tyr Ile Leu Ala 30 165 170 175
 - Ile Thr Ser Phe Leu Glu Phe Val Ile Pro Val Ile Leu Val Ala Tyr 180 185 190
 - Phe Asn Met Asn Ile Tyr Trp Ser Leu Trp Lys Arg Asp His Leu Ser 195 200 205
- Arg Cys Gln Ser His Pro Gly Leu Thr Ala Val Ser Ser Asn Ile Cys 210 215 220

	Gly 225	His	Ser	Phe	Arg	Gly 230	Arg	Leu	Ser	Ser	Arg 235		Ser	Leu	Ser	Al 24
	Ser	Thr	Glu	Val	Pro 245	Ala	Ser	Phe	His	Ser 250		Arg	Gln	Arg	Arg 255	Lу
5	Ser	Ser	Leu	Met 260	Phe	Ser	Ser	Arg	Thr 265	Lys	Met	Asn	Ser	Asn 270	Thr	Il
	Ala	Ser	Lys 275	Met	Gly	Ser	Phe	Ser 280	Gln	Ser	Asp	Ser	Val 285		Leu	His
0	Gln	Arg 290	Glu	His	Val	Glu	Leu 295	Leu	Arg	Ala	Arg	Arg 300	Leu	Ala	Lys	Sei
•	Leu 305	Ala	Ile	Leu	Leu	Gly 310	Val	Phe	Ala	Val	Cys 315	Trp	Ala	Pro	Tyr	Ser 320
٠	Leu	Phe	Thr	Ile	Val 325	Leu	Ser	Phe	Tyr	Ser 330	Ser	Ala	Thr	Gly	Pro 335	Lys
5	Ser	Val	Trp	Tyr 340	Arg	Ile	Ala	Phe	Trp 345	Leu	Gln	Trp	Phe	Asn 350	Ser	Phe
	Val	Asn	Pro 355	Leu	Leu	Tyr		Leu 360	Cys	His	Lys	Arg	Phe 365	Gln	Lys	Àla
)	Phe	Leu 370	Lys	Ile	Phe		Ile 375	Lys	Lys	Gl'n	Pro	Leu 380	Pro	Ser	Gln	His
	Ser 385	Arg	Ser	Val		Ser 390									-	
(16)	INFO	RMAT	NOI	FOR	SEQ	ID N	0:15	:								

(i) SEQUENCE CHARACTERISTICS:

25 (A) LENGTH: 1128 base pairs

- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- 30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

ATGGCGAACG CGAGCGAGCC GGGTGGCAGC GGCGGCGGC AGGCGCCGC CCTGGGCCTC 60

AAGCTGGCCA CGCTCAGCCT GCTGCTGCC GTGAGCCTAG CGGGCAACGT GCTCGCCG 120

CTGCTGATCG TGCGGGAGCG CAGCCTGCAC CGCGCCCGT ACTACCTGCT GCTCGACCTG 180

TGCCTGGCCG ACGGCTGCG CGCGCCCCG GCCCCGG CCGTCATGCT GGCGGCGCG 240

35 CGTGCGGCGG CCGCGC GGCGCCCG GCCGCCGC GCTCGCAAGCT GCTCGCCTTC 300

			*			
CTGGCCGCGC	TCTTCTGCTT	CCACGCCGCC	TTCCTGCTGC	TGGGCGTGG	GTCACCCGC	360
TACCTGGCCA	TCGCGCACCA	CCGCTTCTAT	GCAGAGCGCC	TGGCCGGCTG	GCCGTGCGCC	420
GCCATGCTGG	TGTGCGCCGC	CTGGGCGCTG	GCGCTGGCCG	CGGCCTTCCC	GCCAGTGCTG	480
GACGGCGGTG	GCGACGACGA	GGACGCCCC	TGCGCCCTGG	AGCAGCGGCC	CGACGCCCC	540
5 CCCGGCGCGC	TGGGCTTCCT	GCTGCTGCTG	GCCGTGGTGG	TGGGCGCCAC	GCACCTCGTC	600
TACCTCCGCC	TGCTCTTCTT	CATCCACGAC	CGCCGCAAGA	TGCGGCCCGC	GCGCCTGGTG	660
CCCGCCGTCA	GCCACGACTG	GACCTTCCAC	GGCCCGGGCG	CCACCGGCCA	GGCGGCCGCC	720
AACTGGACGG	CGGGCTTCGG	CCGCGGGCCC	ACGCCGCCCG	CGCTTGTGGG	CATCCGGCCC	780
GCAGGGCCGG	GCCGCGCGC	GCGCCGCCTC	CTCGTGCTGG	AAGAATTCAA	GACGGAGAAG	840
0 AGGCTGTGCA	AGATGTTCTA	CGCCGTCACG	CTGCTCTTCC	TGCTCCTCTG	GGGGCCCTAC	900
GTCGTGGCCA	GCTACCTGCG	GGTCCTGGTG	CGGCCCGGCG	CCGTCCCCCA	GGCCTACCTG	960
ACGGCCTCCG	TGTGGCTGAC	CTTCGCGCAG	GCCGGCATCA	ACCCCGTCGT	GTGCTTCCTC1	020
TTCAACAGGG	AGCTGAGGGA	CTGCTTCAGG	GCCCAGTTCC	CCTGCTGCCA	GAGCCCCCGG1	080
ACCACCCAGG	CGACCCATCC	CTGCGACCTG	AAAGGCATTG	GTTTATGA	1	128

15 (17) INFORMATION FOR SEQ ID NO:16:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 375 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS:
- 20 (D) TOPOLOGY: not relevant
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:
 - Met Ala Asn Ala Ser Glu Pro Gly Gly Ser Gly Gly Glu Ala Ala 1 5 10 15
- 25 Ala Leu Gly Leu Lys Leu Ala Thr Leu Ser Leu Leu Cys Val Ser 20 25 30
 - Leu Ala Gly Asn Val Leu Phe Ala Leu Leu Ile Val Arg Glu Arg Ser 35 40 45
- Leu His Arg Ala Pro Tyr Tyr Leu Leu Leu Asp Leu Cys Leu Ala Asp 50 55 60
 - Gly Leu Arg Ala Leu Ala Cys Leu Pro Ala Val Met Leu Ala Ala Arg
 65 70 75 80

	Arg	Ala	Ala	Ala	Al.a 85	Ala	Gly	Ala	Pro	Pro 90	Gly	Ala	Leu	Gly	Сув 95	Lys ,
	Leu	Leu	Ala	Phe 100		Ala	Ala	Leu	Phe 105		Phe	His	Ala	Ala 110		Leu
5	Leu	Leu	Gly 115		Gly	Val	Thr	Arg 120		Leu	Ala	Ile	Ala 125	His	His	Arg
	Phe	Тут 130	Ala	Glu	Arg	Leu	Ala 135		Trp	Pro	Cys	Ala 140	Ala	Met	Leu	Val
10	Cys 145	Ala	Ala	Trp	Ala	Leu 150		Leu	Ala	Ala	Ala 155		Pro	Pro	Val	Leu 160
	Asp	Gly	.Gly	Gly	Asp 165	Asp	Glu	Asp	Ala	Pro 170	Сув	Ala	Leu	Glu	Gln 175	Arg
	Pro	Asp	Gly	Ala 180	Pro	Gly	Ala	Leu	Gly 185	Phe	Leu	Leu	Leu	Leu 190	Ala	Val
15	Val	Val	Gly 195		Thr	His	Leu	Val 200	Tyr	Leu	Arg	Leu	Leu 205	Phe	Phe	Ile
. •	'His	Asp 210	Arg	Arg	Lys	Met	Arg 215	Pro	Ala	Arg	Leu	Val 220	Pro	Ala	Val	Ser
20	His 225	Asp	Trp	Thr	Phe	His 230	Gly	Pro	Gly	Ala	Thr 235	Gly	Gln	Ala	Ala	Ala 240
	Asņ	Trp	Thr	Ala	Gly 245	Phe	Gly	Arg	Gly	Pro 250	Thr	Pro	Pro	Ala	Leu 255	Val
	Gly	Ile	Arg	Pro 260	Ala	Gly	Pro	Gly	Arg 265	Gly	Ala	Arg	Arg	Leu 270	Leu	Val
25	Leu	Glu	Glu 275	Phe	Lys	Thr	Glu	Lys 280	Arg	Leu	Cys	Lys	Met 285	Phe	Tyr	Ala
- :	.Val	Thr 290	Leu	Leu	Phe	Leu	Leu 295	Leu	Trp	Gly	Pro	Tyr 300	Val	Val	Ala	Ser
30	Tyr 305	Leu	Arg	Val	Leu.	Val 310	Arg	Pro	Gly	Ala	Val 315	Pro	Gln	Ala	Tyr	Leu 320
	Thr	Ala	Ser	Val	Trp 325	Leu	Thr	Phe-	Ala	Gln 330	Ala	Gly	Ile	Asn	Pro 335	Val
	Val	Суз	Phe	Leu 340	Phe	Asn	Arg	Glu	Leu 345	Arg	Asp	Cys		Arg 350	Ala	Gln
35	Phe	Pro	Cys 355	Cys	Gln	Ser	Pro	Arg 360	Thr	Thr	Gln		Thr 365	His	Pro	Cys
	Asp	Leu	Lys	Gly	Ile	Gly	Leu			•	• :					

375

- (18) INFORMATION FOR SEQ ID NO:17:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1002 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA (genomic)
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:
- 10 ATGAACACCA CAGTGATGCA AGGCTTCAAC AGATCTGAGC GGTGCCCCAG AGACACTCGG 60 ATAGTACAGC TGGTATTCCC AGCCCTCTAC ACAGTGGTTT TCTTGACCGG CATCCTGCTG 120 AATACTTTGG CTCTGTGGGT GTTTGTTCAC ATCCCCAGCT CCTCCACCTT CATCATCTAC 180 CTCAAAAACA CTTTGGTGGC CGACTTGATA ATGACACTCA TGCTTCCTTT CAAAATCCTC 240 TCTGACTCAC ACCTGGCACC CTGGCAGCTC AGAGCTTTTG TGTGTCGTTT TTCTTCGGTG 300 15 ATATTTTATG AGACCATGTA TGTGGGCATC GTGCTGTTAG GGCTCATAGC CTTTGACAGA 360 TTCCTCAAGA TCATCAGACC TTTGAGAAAT ATTTTTCTAA AAAAACCTGT TTTTGCAAAA 420 ACGGTCTCAA TCTTCATCTG GTTCTTTTTG TTCTTCATCT CCCTGCCAAA TACGATCTTG 480 AGCAACAAGG AAGCAACACC ATCGTCTGTG AAAAAGTGTG CTTCCTTAAA GGGGCCTCTG 540 GGGCTGAAAT GGCATCAAAT GGTAAATAAC ATATGCCAGT TTATTTTCTG GACTGTTTTT 600 20 ATCCTAATGC TTGTGTTTTA TGTGGTTATT GCAAAAAAAG TATATGATTC TTATAGAAAG 660 TCCAAAAGTA AGGACAGAAA AAACAACAAA AAGCTGGAAG GCAAAGTATT TGTTGTCGTG 720 GCTGTCTTCT TTGTGTGTTT TGCTCCATTT CATTTTGCCA GAGTTCCATA TACTCACAGT 780 CAAACCAACA ATAAGACTGA CTGTAGACTG CAAAATCAAC TGTTTATTGC TAAAGAAACA 840 ACTCTCTTTT TGGCAGCAAC TAACATTTGT ATGGATCCCT TAATATACAT ATTCTTATGT 900 25 AAAAAATTCA CAGAAAAGCT ACCATGTATG CAAGGGAGAA AGACCACAGC ATCAAGCCAA 960 GAAAATCATA GCAGTCAGAC AGACAACATA ACCTTAGGCT GA 1002
 - (19) INFORMATION FOR SEQ ID NO:18:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 333 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS:

(D)	TOPOLOGY:	not	relevant

(ii) MOLECULE TYPE: protein

	(xi)	SEQ	UENC	E DE	SCRI	PTIC	N: S	EQ I	D NC	:18:						
5	Met 1	Asn	Thr	Thr	Val 5	Met	Gln	Gly	Phe	Asn 10	Arg	Ser	Glu	Årg	Cys 15	Pro
	Arg	Asp	Thr	Arg 20	Ile	Val	Gln	Leu	Val 25	Phe	Pro	Ala	Leu	Туr 30	Thr	Val
	Val	Phe	Leu 35	Thr	Gly	Ile	Leu	Leu 40	Asn	Thr	Leu	Ala	Leu 45	Trp	Val	Phe
10	Val	His 50	Ile	Pro	Ser	Ser	Ser 55	Thr	Phe	Ile	Ile	Tyr 60	Leu	Lys	Asn	Thŗ
	Leu 65	Val	Ala	Asp	Leu	Ile 70	Met	Thr	Leu	Met	Leu 75	Pro	Phe	Lys	·Ile	Leu 80
15	Ser	Asp	Ser	His	Leú 85	Ala	Pro	Trp	Gln	Leu 90	Arg	Ala	Phe	Val	Cys 95	Arg
	Phe	Ser	Ser	Val 100	Ile	Phe	Tyr	Glu	Thr 105		Tyr	Val	Gly	Ile 110	Val	Leu
	Leu	Gly	Leu 115	Ile	Ala	Phe	Asp	Arg 120	Phe	Leu	Lys	Ile	Ile 125	Arg	Pro	Leu
20	Arg	Asn 130	Ile	Phe	Leu	Lys	Lys 135	Pro	Val	Phe	Ala	Lys 140	Thr	Val	Ser	Ile
	Phe 145	Ile	Trp	Phe	Phe	Leu 150	Phe	Phe	Ile	Ser	Leu 155	Pro	Asn.	Thr	Ile	Leu 160
25	Ser	Asn	Lys	Glu	Ala 165	Thr	Pro	Ser	Ser	Val 170	Lys	Lys	Cys	Ala	Ser 175	
	Lys	Gly	Pro	Leu 180	Gly	Leu	Lys	Trp	His 185	Gln	Met	Val	Asn	Asn 190	Ile	Cys
	Gln	Phe	Ile 195	Phe	Trp	Thr	Val	Phe 200	Ile	Leu	Met	Leu	Val 205		Tyr	Val
30	Val	Ile 210	Ala	Lys	Lys	Val	Tyr 215	Asp	Ser	Tyr	Arg	Lys 220	Ser	Lys	Ser	Lys
	Asp 225	Arg	Lys	Asn		Lys 230	Lys	Leu	Glu		Lys 235	Val	Phe	Vaİ	Val	Val 240
35	Ala	Val	Phe		Val 245		Phe	Ala		Phe 250	His	Phe	Ala	Arg	Val 255	Pro

- Tyr Thr His Ser Gln Thr Asn Asn Lys Thr Asp Cys Arg Leu Gln Asn 260 265 270
- Gln Leu Phe Ile Ala Lys Glu Thr Thr Leu Phe Leu Ala Ala Thr Asn 275 280 285
- 5 Ile Cys Met Asp Pro Leu Ile Tyr Ile Phe Leu Cys Lys Lys Phe Thr 290 295 300
 - Glu Lys Leu Pro Cys Met Gln Gly Arg Lys Thr Thr Ala Ser Ser Gln 305 310 315 320
- Glu Asn His Ser Ser Gln Thr Asp Asn Ile Thr Leu Gly
 325 330
 - (20) INFORMATION FOR SEQ ID NO:19:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1122 base pairs
 - (B) TYPE: nucleic acid
- 15 (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA (genomic)
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:
- ATGGCCAACA CTACCGGAGA GCCTGAGGAG GTGAGCGGCG CTCTGTCCC ACCGTCCCA 600

 20 TCAGCTTATG TGAAGCTGGT ACTGCTGGGA CTGATTATGT GCGTGAGCCT GGCGGGTAAC 120

 GCCATCTTGT CCCTGCTGG GCTCAAGGAG CGTGCCCTGC ACAAGGCTCC TTACTACTTC 180

 CTGCTGGACC TGTGCCTGGC CGATGGCATA CGCTCTGCCG TCTGCTTCCC CTTTGTGCTG 240

 GCTTCTGTGC GCCACGGCTC TTCATGGACC TTCAGTGCAC TCAGCTGCAA GATTGTGGCC 300

 TTTATGGCCG TGCTCTTTG CTTCCATGCG GCCTTCATGC TGTTCTGCAT CAGCGTCACC 360

 25 CGCTACATGG CCATCGCCCA CCACCGCTTC TACGCCAAGC GCATGACACT CTGGACATGC 420

 GCGGCTGTCA TCTGCATGGC CTGGACCCTG TCTGTGGCCA TGGCCTTCCC ACCTGTTT 480

 GACGTGGGCA ATGACACGCT GGGCTTCATG CTTATGTTGG CTGTCCTAT GCAGCTACC 540

 TTCAAGGCCA ATGACACGCT GGCCTTCATG CTTATGTTGG CTGTGCTCAT GACACTTCC 600

 CATGCTGTCT ACGGCAAGCT GCTCCTCTC GAGTATCGTC ACCGCAAGAT GAAGCCAGTG 660

 30 CAGATGGTGC CAGCCATCAG CCAGAACTGG ACATTCCATG GTCCCGGGGC CACCGGCCAG 720

 GCTGCTGCCA ACTGGATCGC CGGCTTTGGC CGTGGGCCCA TGCCACCAAC CCTGCTGGGT 780

ATCCGGCAGA ATGGGCATGC AGCCAGCCGG CGGCTACTGG GCATGGACGA GGTCAAGGGT 840

GAAAAGCAGC TGGGCCGCAT GTTCTACGCG ATCACACTGC TCTTTCTGCT CCTCTGGTCA 900

CCCTACATCG TGGCCTGCTA CTGGCGAGTG TTTGTGAAAG CCTGTGCTGT GCCCCACCGC 960

TACCTGGCCA CTGCTGTTTG GATGAGCTTC GCCCAGGCTG CCGTCAACCC AATTGTCTGC1020

TTCCTGCTCA ACAAGGACCT CAAGAAGTGC CTGACCACTC ACGCCCCTG CTGGGGCACA1080

5 GGAGGTGCCC CGGCTCCCAG AGAACCCTAC TGTGTCATGT GA 1122

- (21) INFORMATION FOR SEQ ID NO:20:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 373 amino acids
 - (B) TYPE: amino acid
- 10
- (C) STRANDEDNESS:
- (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:
- Met Ala Asn Thr Thr Gly Glu Pro Glu Glu Val Ser Gly Ala Leu Ser
 15 1 5 10 15
 - Pro Pro Ser Ala Ser Ala Tyr Val Lys Leu Val Leu Leu Gly Leu Ile 20 25 30
 - Met Cys Val Ser Leu Ala Gly Asn Ala Ile Leu Ser Leu Leu Val Leu 35 40 45
- 20 Lys Glu Arg Ala Leu His Lys Ala Pro Tyr Tyr Phe Leu Leu Asp Leu
 50 55
 - Cys Leu Ala Asp Gly Ile Arg Ser Ala Val Cys Phe Pro Phe Val Leu 65 70 75 80
- Ala Ser Val Arg His Gly Ser Ser Trp Thr Phe Ser Ala Leu Ser Cys

 85 90 95
 - Lys Ile Val Ala Phe Met Ala Val Leu Phe Cys Phe His Ala Ala Phe 100 105 110
 - Met Leu Phe Cys Ile Ser Val Thr Arg Tyr Met Ala Ile Ala His His 115 120 125
- Arg Phe Tyr Ala Lys Arg Met Thr Leu Trp Thr Cys Ala Ala Val Ile 130 135 140
 - Cys Met Ala Trp Thr Leu Ser Val Ala Met Ala Phe Pro Pro Val Phe 145 150 155 160
- Asp Val Gly Thr Tyr Lys Phe Ile Arg Glu Glu Asp Gln Cys Ile Phe 165 170 175

- Glu His Arg Tyr Phe Lys Ala Asn Asp Thr Leu Gly Phe Met Leu Met 185 Leu Ala Val Leu Met Ala Ala Thr His Ala Val Tyr Gly Lys Leu Leu Leu Phe Glu Tyr Arg His Arg Lys Met Lys Pro Val Gln Met Val Pro Ala Ile Ser Gln Asn Trp Thr Phe His Gly Pro Gly Ala Thr Gly Gln Ala Ala Ala Asn Trp Ile Ala Gly Phe Gly Arg Gly Pro Met Pro Pro 10 Thr Leu Leu Gly Ile Arg Gln Asn Gly His Ala Ala Ser Arg Arg Leu Leu Gly Met Asp Glu Val Lys Gly Glu Lys Gln Leu Gly Arg Met Phe 280 Tyr Ala Ile Thr Leu Leu Phe Leu Leu Leu Trp Ser Pro Tyr Ile Val 15 Ala Cys Tyr Trp Arg Val Phe Val Lys Ala Cys Ala Val Pro His Arg Tyr Leu Ala Thr Ala Val Trp Met Ser Phe Ala Gln Ala Ala Val Asn 20 Pro Ile Val Cys Phe Leu Leu Asn Lys Asp Leu Lys Lys Cys Leu Thr 340 Thr His Ala Pro Cys Trp Gly Thr Gly Gly Ala Pro Ala Pro Arg Glu 360

30

Pro Tyr Cys Val Met

(22) INFORMATION FOR SEQ ID NO:21:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1053 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:
- 35 ATGGCTTTGG AACAGAACCA GTCAACAGAT TATTATTATG AGGAAAATGA AATGAATGGC 60
 ACTTATGACT ACAGTCAATA TGAATTGATC TGTATCAAAG AAGATGTCAG AGAATTTGCA 120

AAAGTTTTCC TCCCTGTATT CCTCACAATA GCTTTCGTCA TTGGACTTGC AGGCAATTCC 180 ATGGTAGTGG CAATTTATGC CTATTACAAG AAACAGAGAA CCAAAACAGA TGTGTACATC 240 CTGAATTTGG CTGTAGCAGA TTTACTCCTT CTATTCACTC TGCCTTTTTG GGCTGTTAAT 300 GCAGTTCATG GGTGGGTTTT AGGGAAAATA ATGTGCAAAA TAACTTCAGC CTTGTACACA 360 5 CTAAACTTTG TCTCTGGAAT GCAGTTTCTG GCTTGCATCA GCATAGACAG ATATGTGGCA 420 GTAACTAATG TCCCCAGCCA ATCAGGAGTG GGAAAACCAT GCTGGATCAT CTGTTTCTGT 480 GTCTGGATGG CTGCCATCTT GCTGAGCATA CCCCAGCTGG TTTTTTATAC AGTAAATGAC 540 AATGCTAGGT GCATTCCCAT TTTCCCCCGC TACCTAGGAA CATCAATGAA AGCATTGATT 600 CAAATGCTAG AGATCTGCAT TGGATTTGTA GTACCCTTTC TTATTATGGG GGTGTGCTAC 660 10 TTTATCACGG CAAGGACACT CATGAAGATG CCAAACATTA AAATATCTCG ACCCCTAAAA 720 GTTCTGCTCA CAGTCGTTAT AGTTTTCATT GTCACTCAAC TGCCTTATAA CATTGTCAAG 780 TTCTGCCGAG CCATAGACAT CATCTACTCC CTGATCACCA GCTGCAACAT GAGCAAACGC 840 ATGGACATEG CCATCCAAGT CACAGAAAGC ATTGCACTCT TTCACAGCTG CCTCAACCCA 900 ATCCTTTATG TTTTTATGGG AGCATCTTTC AAAAACTACG TTATGAAAGT GGCCAAGAAA 960 15 TATGGGTCCT GGAGAAGACA GAGACAAAGT GTGGAGGAGT TTCCTTTTGA TTCTGAGGGT1020 CCTACAGAGC CAACCAGTAC TTTTAGCATT TAA 1053

(23) INFORMATION FOR SEQ ID NO:22:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 350 amino acids
- 20 (B) TYPE: amino acid
 - (C) STRANDEDNESS:
 - (D) TOPOLOGY: not relevant
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:
- 25 Met Ala Leu Glu Gln Asn Gln Ser Thr Asp Tyr Tyr Glu Glu Asn 1 5 10
 - Glu Met Asn Gly Thr Tyr Asp Tyr Ser Gln Tyr Glu Leu Ile Cys Ile
 20 25 30
- Lys Glu Asp Val Arg Glu Phe Ala Lys Val Phe Leu Pro Val Phe Leu 30 35 40 45

Thr Ile Ala Phe Val Ile Gly Leu Ala Gly Asn Ser Met Val Val Ala

		50	•				55					60	٠.			
	I10 65	е Ту	r Al	а Ту:	г Ту	r Ly: 70	s Ly	s Gl:	n Ar	g Th	r Ly: 75	s Th	r As	p Va	l Ty	r Il 80
5	Let	ı Ası	n Le	u Ala	a Val 85	l Ala	a Ası	Le	u Lei	u Lei 90	u Let	ı Phe	e Thi	r Lei	1 Pro	Ph
	Tr	Ala	a Va	l Ası 100	1 Ala	a Val	l His	Gl;	7 Trp 109	p Vai	l Lei	ı Gly	y Lys	11¢		: Су
			. 11:	٠.	:			120)				125	5		•
10		130	,	a Cys			135	i		·		140				
	1.45	•		ı Ser		150	۲				155	٠			•	160
15			•	Ala	165					170	٠				175	
•				Asp 180					185					190	•	
			195					200					205			
20		210		Pro			215	,				220				
	225			Met		230					235	,				240
25				Thr	245	٠				250					255	
				Lys 260	*				265		٠.			270		•
			275	Asn				280					285			
30		290		Ala			295		•			300				
	305		•	Ala		310					315				. ,	320
35		•		Trp	325					.330	٠.		*		Pro 335	Phe
	Asp	Ser	Glu	Gly 340	Pro	Thr	Glu	Pro	Thr 345	Ser	Thr	Pḥe	Ser	Ile 350		

(24) INFORMATION FOR SEQ ID NO:23:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1116 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:
- ATGCCAGGAA ACGCCACCC AGTGACCACC ACTGCCCCGT GGGCCTCCCT GGGCCTCTCC 60 10 GCCAAGACCT GCAACAACGT GTCCTTCGAA GAGAGCAGGA TAGTCCTGGT CGTGGTGTAC 120 AGCGCGGTGT GCACGCTGGG GGTGCCGGCC AACTGCCTGA CTGCGTGGCT GGCGCTGCTG 180 CAGGTACTGC AGGGCAACGT GCTGGCCGTC TACCTGCTCT GCCTGGCACT CTGCGAACTG 240 CTGTACACAG GCACGCTGCC ACTCTGGGTC ATCTATATCC GCAACCAGCA CCGCTGGACC 300 CTAGGCCTGC TGGCCTCGAA GGTGACCGCC TACATCTTCT TCTGCAACAT CTACGTCAGC 360 15 ATCCTCTTCC TGTGCTGCAT CTCCTGCGAC CGCTTCGTGG CCGTGGTGTA CGCGCTGGAG 420 AGTCGGGGCC GCCGCCGC GAGGACCGCC ATCCTCATCT CCGCCTGCAT CTTCATCCTC 480 GTCGGGATCG TTCACTACCC GGTGTTCCAG ACGGAAGACA AGGAGACCTG CTTTGACATG 540 CTGCAGATGG ACAGCAGGAT TGCCGGGTAC TACTACGCCA GGTTCACCGT TGGCTTTGCC 600 ATCCCTCTCT CCATCATCGC CTTCACCAAC CACCGGATTT TCAGGAGCAT CAAGCAGAGC 660 20 ATGGGCTTAA GCGCTGCCCA GAAGGCCAAG GTGAAGCACT CGGCCATCGC GGTGGTTGTC 720 ATCTTCCTAG TCTGCTTCGC CCCGTACCAC CTGGTTCTCC TCGTCAAAGC CGCTGCCTTT 780 TCCTACTACA GAGGAGACAG GAACGCCATG TGCGGCTTGG AGGAAAGGCT GTACACAGCC 840 TCTGTGGTGT TTCTGTGCCT GTCCACGGTG AACGGCGTGG CTGACCCCAT TATCTACGTG 900 CTGGCCACGG ACCATTCCCG CCAAGAAGTG TCCAGAATCC ATAAGGGGTG GAAAGAGTGG 960 25 TCCATGAAGA CAGACGTCAC CAGGCTCACC CACAGCAGGG ACACCGAGGA GCTGCAGTCG1020 CCCGTGGCCC TTGCAGACCA CTACACCTTC TCCAGGCCCG TGCACCCACC AGGGTCACCA1080 TGCCCTGCAA AGAGGCTGAT TGAGGAGTCC TGCTGA 1116
 - (25) INFORMATION FOR SEQ ID NO:24:
 - (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 371 amino acids

(B)	TYPE:	amino	2014

- (C) STRANDEDNESS:
- (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: protein
- 5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:
 - Met Pro Gly Asn Ala Thr Pro Val Thr Thr Ala Pro Trp Ala Ser

 10 15
 - Leu Gly Leu Ser Ala Lys Thr Cys Asn Asn Val Ser Phe Glu Glu Ser 20 25 30
- 10 Arg Ile Val Leu Val Val Val Tyr Ser Ala Val Cys Thr Leu Gly Val 35 40 45
 - Pro Ala Asn Cys Leu Thr Ala Trp Leu Ala Leu Leu Gln Val Leu Gln 50 55 60
- Gly Asn Val Leu Ala Val Tyr Leu Leu Cys Leu Ala Leu Cys Glu Leu
 15 65 70 75 80
 - Leu Tyr Thr Gly Thr Leu Pro Leu Trp Val Ile Tyr Ile Arg Asn Gln 85 90 95
 - His Arg Trp Thr Leu Gly Leu Leu Ala Ser Lys Val Thr Ala Tyr Ile 100 105 110
- 20 Phe Phe Cys Asn Ile Tyr Val Ser Ile Leu Phe Leu Cys Cys Ile Ser 115 120 125
 - Cys Asp Arg Phe Val Ala Val Val Tyr Ala Leu Glu Ser Arg Gly Arg 130 135 140
- Arg Arg Arg Thr Ala Ile Leu Ile Ser Ala Cys Ile Phe Ile Leu 25 145 150 155 160
 - Val Gly Ile Val His Tyr Pro Val Phe Gln Thr Glu Asp Lys Glu Thr
 165 170 175
 - Cys Phe Asp Met Leu Gln Met Asp Ser Arg Ile Ala Gly Tyr Tyr Tyr 180 185 190
- Ala Arg Phe Thr Val Gly Phe Ala Ile Pro Leu Ser Ile Ile Ala Phe
 195 200 205
 - Thr Asn His Arg Ile Phe Arg Ser Ile Lys Gln Ser Met Gly Leu Ser 210 215 220
- Ala Ala Gln Lys Ala Lys Val Lys His Ser Ala Ile Ala Val Val
 35 225 230 235 240

- Ile Phe Leu Val Cys Phe Ala Pro Tyr His Leu Val Leu Leu Val Lys
 245 250 255
- Ala Ala Ala Phe Ser Tyr Tyr Arg Gly Asp Arg Asn Ala Met Cys Gly
 260 265 270
- 5 Leu Glu Glu Arg Leu Tyr Thr Ala Ser Val Val Phe Leu Cys Leu Ser 275 280 285
 - Thr Val Asn Gly Val Ala Asp Pro Ile Ile Tyr Val Leu Ala Thr Asp 290 295 300
- His Ser Arg Gln Glu Val Ser Arg Ile His Lys Gly Trp Lys Glu Trp 10 305 310 315 320
 - Ser Met Lys Thr Asp Val Thr Arg Leu Thr His Ser Arg Asp Thr Glu 325 330 335
 - Glu Leu Gln Ser Pro Val Ala Leu Ala Asp His Tyr Thr Phe Ser Arg 340 345 350
- Pro Val His Pro Pro Gly Ser Pro Cys Pro Ala Lys Arg Leu Ile Glu 355 360 365

Glu Ser Cys 370

(26) INFORMATION FOR SEQ ID NO:25:

- 20 (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1113 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- 25 (ii) MOLECULE TYPE: DNA (genomic)
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:
- ATGGCGAACT ATAGCCATGC AGCTGACAAC ATTTTGCAAA ATCTCTCGCC TCTAACAGCC 60
 TTTCTGAAAC TGACTTCCTT GGGTTTCATA ATAGGAGTCA GCGTGGTGGG CAACCTCCTG 120
 ATCTCCATTT TGCTAGTGAA AGATAAGACC TTGCATAGAG CACCTTACTA CTTCCTGTTG 180
 30 GATCTTTGCT GTTCAGATAT CCTCAGATCT GCAATTTGTT TCCCATTTGT GTTCAACTCT 240
 GTCAAAAATG GCTCTACCTG GACTTATGGG ACTCTGACTT GCAAAGTGAT TGCCTTTCTG 300
 GGGGTTTTGT CCTGTTTCCA CACTGCTTTC ATGCTCTTCT GCATCAGTGT CACCAGATAC 360
 TTAGCTATCG CCCATCACCG CTTCTATACA AAGAGGCTGA CCTTTTGGAC GTGTCTGGCT 420

GTGATCTGTA TGGTGTGGAC TCTGTCTGTG GCCATGGCAT TTCCCCCGGT TTTAGACGTG 480

GGCACTTACT CATTCATTAG GGAGGAAGAT CAATGCACCT TCCAACACCG CTCCTTCAGG 540
GCTAATGATT CCTTAGGATT TATGCTGCTT CTTGCTCTCA TCCTCCTAGC CACACAGCTT 600
GTCTACCTCA AGCTGATATT TTTCGTCCAC GATCGAAGAA AAATGAAGCC AGTCCAGTTT 660
GTAGCAGCAG TCAGCCAGAA CTGGACTTTT CATGGTCCT GAGCCAGTGG CCAGGCCAGCT 720
5GCCAATTGGC TAGCAGGATT TGGAAGGGGT CCCACACCAC CCACCTTGCT GGGCATCAGG 780
CAAAAATGCAA ACACCACAGG CAGAAGAAGA CTTTTCTGT TAGACCAGTT CAAAAATGGAG 840
AAAAGAATCA GCAGAATGTT CTATATAATG ACTTTTCTGT TTCTAACCTT GTGGGGCCCC 900
TACCTGGTGG CCTGTTATTG GAGAGTTTTT GCCAAGAGGGC CTGTAGTACC AGGGGGATTT 960
CTAACAGCTG CTGTCTGGAT GAGTTTTGCC CAAGCAGCAC TCATTCCTT TGTCTGCATT1020
10 TTCTCAAACA GGGAGCTGAG GCGCTGTTTC AGCACAACCC TTCTTTACTG CAGAAAAATCC1080
AGGTTACCAA GGGAACCTTA CTGTGTTATA TGA

- (27) INFORMATION FOR SEQ ID NO:26:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 370 amino acids
 - (B) TYPE: amino acid

- (C) STRANDEDNESS:
- (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: protein
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:
- Met Ala Asn Tyr Ser His Ala Ala Asp Asn Ile Leu Gln Asn Leu Ser

 1 5 10 15
 - Pro Leu Thr Ala Phe Leu Lys Leu Thr Ser Leu Gly Phe Ile Ile Gly 20 25 30
- Val Ser Val Val Gly Asn Leu Leu Ile Ser Ile Leu Leu Val Lys Asp

 35 40 45
 - Lys Thr Leu His Arg Ala Pro Tyr Tyr Phe Leu Leu Asp Leu Cys Cys 50 55 60
 - Ser Asp Ile Leu Arg Ser Ala Ile Cys Phe Pro Phe Val Phe Asn Ser 65 70 75 80
- Val Lys Asn Gly Ser Thr Trp Thr Tyr Gly Thr Leu Thr Cys Lys Val
 - Ile Ala Phe Leu Gly Val Leu Ser Cys Phe His Thr Ala Phe Met Leu

				100					105	5 ,				110)	
	Phe	cys	115	Ser	Val	Thr	Arg	Tyr 120		. Ala	Ile	Ala	His 125		Arg	Phe
5	Туг	Thr 130	Lys	Arg	Leu	Thr	Phe 135		Thr	Cys	Leu	Ala 140		Ile	Cys	Met
	Val 145	Trp	Thr	Leu	Ser	Val 150		Met	Ala	Phe	Pro 155		Val	Leu	Asp	Val
	Gly	Thr	Tyr	Ser	Phe 165	Ile	Arg	Glu	Glu	Asp 170	Gln	Cys	Thr	Phe	Gln 175	His
10	Arg	Ser	Phe	Arg 180	Ala	Asn	Asp	Ser	Leu 185		Phe	Met	Leu	Leu 190		Ala
	Leu	Ile	Leu 195	Leu	Ala	Thr	Gln	Leu 200	Val	Tyr	Leu	Lys	Leu 205	Ile	Phe	Phe
15	Val	His 210	Asp	Arg	Arg	Lys	Met 215	ГÀЗ	Pro	Val		Phe 220	Val	Ala	Ala	Val
	Ser 225	Gln	Asn	Trp	Thr	Phe 230	His	Gly	Pro	Gly	Ala 235	Ser	Gly	Gln	Ala	Ala 240
	Ala	Asn	Trp	Leu	Ala 245	Gly	Phe	Gly	Arg	Gly 250	Pro	Thr	Pro	Pro	Thr 255	Leu
20	Leu	Gly	Ile	Arg 260	Gln	Asn	Ala	Asn	Thr 265	Thr	Gly	Arg	Arg	Arg 270	Leu	Leu
٠	Val	Leu	Asp 275	Glu	Phe	Lys	Met	Glu 280	Lys	Arg	Ile	Ser	Arg 285	Met	Phe	Tyr
25	Ile	Met 290	Thr	Phe	Leu	Phe	Leu 295	Thr	Leu	Trp	Gly	Pro 300	Tyr	Leu	Val	Ala
	Cys 305	Tyr	Trp	Arg		Phe 310	Ala	Arg	Gly	Pro	Val 315	Val	Pro	Gly	Gly	Phe 320
	Leu	Thr	Ala	Ala	Val 325	Trp	Met	Ser	Phe	Ala 330	Gln	Ala	Gly	Ile	Asn 335	Pro
30	Phe	Val	Cys	Ile 340	Phe	Ser	Asn	Arg	Glu 345	Leu	Arg	Arg	Cys	Phe 350	Ser-	Thr
. ,	Thr	Leu	Leu 355	Tyr	Cys	Arg		Ser 360	Arg	Leu	Pro	Arg	Glu 365	Pro	Tyr	.Cys.
35	Val	Ile 370														
(28)	TNEC	יי ליאכו	TOM	EOD	ORO	TD X										

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1080 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:
- ATGCAGGTCC CGAACAGCAC CGGCCCGGAC AACGCGACGC TGCAGATGCT GCGGAACCCG 60 GCGATCGCGG TGGCCCTGCC CGTGGTGTAC TCGCTGGTGG CGGCGGTCAG CATCCCGGGC 120 10 AACCTCTTCT CTCTGTGGGT GCTGTGCCGG CGCATGGGGC CCAGATCCCC GTCGGTCATC 180 TTCATGATCA ACCTGAGCGT CACGGACCTG ATGCTGGCCA GCGTGTTGCC TTTCCAAATC 240 TACTACCATT GCAACCGCCA CCACTGGGTA TTCGGGGTGC TGCTTTGCAA CGTGGTGACC 300 GTGGCCTTTT ACGCAAACAT GTATTCCAGC ATCCTCACCA TGACCTGTAT CAGCGTGGAG 360 CGCTTCCTGG GGGTCCTGTA CCCGCTCAGC TCCAAGCGCT GGCGCCGCCG TCGTTACGCG 420 15 GTGGCCGCGT GTGCAGGGAC CTGGCTGCTG CTCCTGACCG CCCTGTGCCC GCTGGCGCGC 480 ACCGATCTCA CCTACCCGGT GCACGCCCTG GGCATCATCA CCTGCTTCGA CGTCCTCAAG 540 TGGACGATGC TCCCCAGCGT GGCCATGTGG GCCGTGTTCC TCTTCACCAT CTTCATCCTG 600 CTGTTCCTCA TCCCGTTCGT GATCACCGTG GCTTGTTACA CGGCCACCAT CCTCAAGCTG 660 TTGCGCACGG AGGAGGCGCA CGGCCGGGAG CAGCGGAGGC GCGCGGTGGG CCTGGCCGCG 720 20 GTGGTCTTGC TGGCCTTTGT CACCTGCTTC GCCCCCAACA ACTTCGTGCT CCTGGCGCAC 780 ATCGTGAGCC GCCTGTTCTA CGGCAAGAGC TACTACCACG TGTACAAGCT CACGCTGTGT 840 CTCAGCTGCC TCAACAACTG TCTGGACCCG TTTGTTTATT ACTTTGCGTC CCGGGAATTC 900 CAGCTGCGCC TGCGGGAATA TTTGGGCTGC CGCCGGGTGC CCAGAGACAC CCTGGACACG 960 CGCCGCGAGA GCCTCTTCTC CGCCAGGACC ACGTCCGTGC GCTCCGAGGC CGGTGCGCAC1020 25 CCTGAAGGGA TGGAGGAGC CACCAGGCCC GGCCTCCAGA GGCAGGAGAG TGTGTTCTGA1080
 - (29) INFORMATION FOR SEQ ID NO:28:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 359 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS:

(D) TOPOLOGY: not relevant

(ii)	MOLECULE	TYPE:	protein
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	(xi) SE	QUENCE	DESCR	IPTIC	DM: 2	SEQ :	ID NO):28:	:				, '	٠.
	Met Gli	n Val	Pro Asi 5	n Sei	Thi	Gly	/ Pro	Asp 10	Asr	ı Ala	Thr	Leu •	Gln 15	. Met
5	Leu Aro	g Asn :	Pro Ala 20	a Ile	a Ala	Val	Ala 25	Let	Pro	Val	. Val	. Tyr 30	Ser	Leu
÷	Val Ala	Ala 35	Val Sei	r Ile	Pro	Gly 40	Asn	Leu	Phe	Ser	Leu 45	Trp	Val	Leu
10	Cys Arg 50	Arg N	Met Gly	/ Pro	Arg 55	Ser	Pro	Ser	Val	Ile 60	Phe	Met	Ile	Asn
	Leu Ser 65	Val 1	Chr Asr	70	Met	Leu	Ala	Ser	Val 75	Leu	Pro	Phe	Gln	Ile 80
	Tyr Tyr	His (Cys Asn 85	Arg	His	His	Trp	Val 90	Phe	Gly	Val	Leu	Leu 95	Cys
15	Asn Val	Val 1	hr Val	Ala	Phe	Tyr	Ala 105	Asn	Met	Tyr	Ser	Ser 110	Ile	Leu
	Thr Met	Thr C	ys Ile	Ser	Val	Glu 120	Arg	Phe	Leu	Gly	Val 125	Leu	Tyr	Pro
20	Leu Ser 130	Ser L	ys Arg	Trp	Arg 135	Arg	Arg	Arg	Tyr	Ala 140	Val	Ala	Ala	Cys
	Ala Gly 145	Thr T	rp Leu	Leu 150	Leu	Leu	Thr	Ala	Leu 155	Cys	Pro	Leu	Ala	Arg 160
	Thr Asp	Leu T	hr Tyr 165	Pro	Val	His	Ala	Leu 170	Gly	Ile	Ile	Thr	Cys 175	Phe
25	Asp Val	Leu L	ys Trp 80	Thr	Met	Leu	Pro 185	Ser	Val	Ala	Met	Trp 190	Ala	Val
	Phe Leu	Phe Ti	hr Ile	Phe	Ile	Leu 200	Leu	Phe	Leu		Pro 205	Phe	Val	Ile
30	Thr Val 210	Ala Cy	ys Tyr		Ala 215	Thr	Ile	Leu		Leu 220	Leu	Arg	Thr	Glu
	Glu Ala 225	His Gi	ly Arg	Glu 230	Gln	Arg	Arg .		Ala 235	Val	Gly	Leu .		Ala 240
	Val Val	Leu Le	245	Phe	Val	Thr		Phe . 250		Pro .	Asn .		Phe '	Val
35	Leu Leu	Ala Hi 26	s Ile	Val :	Ser :		Leu : 265	Phe '	Tyr (Gly :		Ser :	Tyr '	ryr

- His Val Tyr Lys Leu Thr Leu Cys Leu Ser Cys Leu Asn Asn Cys Leu 275 280 285
- Asp Pro Phe Val Tyr Tyr Phe Ala Ser Arg Glu Phe Gln Leu Arg Leu 290 295 300
- Arg Glu Tyr Leu Gly Cys Arg Arg Val Pro Arg Asp Thr Leu Asp Thr 305 310 315 320
 - Arg Arg Glu Ser Leu Phe Ser Ala Arg Thr Thr Ser Val Arg Ser Glu 325 330 335
- Ala Gly Ala His Pro Glu Gly Met Glu Gly Ala Thr Arg Pro Gly Leu
 340 345 350

Gln Arg Gln Glu Ser Val Phe 355

- (30) INFORMATION FOR SEQ ID NO:29:
 - (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1503 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- 20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:
- ATGGAGCGTC CCTGGGAGGA CAGCCCAGGC CCGGAGGGGG CAGCTGAGGG CTCGCCTGTG 60

 CCAGTCGCCG CCGGGGCGC CTCCGGTGCC GCGCCAGTG GCACCAGGCTG GCAGCCATGG 120

 GCTGAGTGCC CGGGACCCAA GGGGAGGGG CAACTGCTGG CGACCGCCGG CCCTTTGCGT 180

 CGCTGGCCCG CCCCCTCGCC TGCCAGCTCC AGCCCCGCCC CCGGAGCGGC GTCCGCTCAC 240

 25 TCGGTTCAAG GCAGCGCGAC TGCGGGTGGC GCACCGCCAG GGCGCAGACC TTGGGGCGCG 300

 CGGCCCATGG AGTCGGGGCT GCTGCGGCCG GCGCCGGTGA GCGAGGTCAT CGTCCTGCAT 360

 TACAACTACA CCGGCAAGCT CCGCGGTGCG AGCTACCAGC CGGGTGCCGG CCTGCGCCC 420

 GACGCCGTGG TGTGCCTGGC GGTGTGCGCC TTCATCGTGC TAGAGAATCT AGCCGTGTTG 480

 TTGGTGCTCG GACGCCACC GCGCTTCCAC GCTCCCATGT TCCTGCTCCT GGGCAGCCT 540

 30 ACGTTGTCGG ATCTGCTGC AGGCGCCGC TACGCCCCA ACATCCTACT GTCGGGGCCG 600

 CTCACCTGCAT AACTGCCCC CGCGCTCTGG TTCGCACGGG AGGGAGGCGT CTTCGTGGCA 660

 CTCACTGCGT CCGTGCTGAC CCTCCTGGC ATCGCCCTGG AGCGCAGCCT CACCATGGCG 720

CGCAGGGGGC CCGCCCGT CTCCAGTCGG GGGCGCACGC TGGCGATGGC AGCCGCGCCC 780
TGGGGCGTGT CGCTGCTCCT CGGGCTCCTG CCAGCGCTGG GCTGGAATTG CCTGGGTCGC 840
CTGGACGCTT GCTCCACTGT CTTGCCGCTC TACGCCAAGG CCTACGTGCT CTTCTGCGTG 900
CTCGCCTTCG TGGGCATCCT GGCCGCGACT TGTGCACTCT ACGCGCGCAT CTACTGCCAG 960
5 GTACGCGCA ACGCGGGGC CCTGCCGGCA CGGCCCGGGA CTGCGGGGAC CACCTCGACC1020
CGGGCGCGCT GCAAGCCGC CTCTCTGGCC TTGCTGCGCA CGCTCAGCGT GGTGCTCCTG1080
GCCTTTGTGG CATGTTGGGG CCCCCTCTTC CTGCTGCTGT TGCTCGACGT GGCGTGCCCG1140
GCGCGCACCT GTCCTGTACT CCTGCAGGCC GATCCCTTCC TGGGACTGGC CATGGCCAAC1200
TCACTTCTGA ACCCCATCAT CTACACGCTC ACCAACCGCG ACCTGCGCA CGCGCTCCTG1260
10 CGCCTGGTCT GCTGCGGACG CCACTCCTCC GGCAGAGACC CGAGTGGCTC CCAGCAGTCG1320
GCGAGCGCGG CTGAGGCTTC CGGGGGCCTG CGCCGCTGCC TGCCCCCGGG CCTTGATGGG1380
AGCTTCAGCG GCTCGGAGCG CTCATCGCCC CAGCGCGACG GGCTGCACAC1500
ACAGGCAGCC CCGGTGCACC CACAGCCGC CGGGCTCCT1440
ACAGGCAGCC CCGGTGCACC CACAGCCGCC CGGGCTCCT1440
ACAGGCAGCC CCGGTGCACC CACAGCCGCC CGGACTCTGG TATCAGAACC GGCTGCAGAC1500

15 (31) INFORMATION FOR SEQ ID NO:30:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 500 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS:
- (D) TOPOLOGY: not relevant
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:
 - Met Glu Arg Pro Trp Glu Asp Ser Pro Gly Pro Glu Gly Ala Ala Glu

 10 15
- 25 Gly Ser Pro Val Pro Val Ala Ala Gly Ala Arg Ser Gly Ala Ala Ala 20 25 30
 - Ser Gly Thr Gly Trp Gln Pro Trp Ala Glu Cys Pro Gly Pro Lys Gly
 35 40 45
- Arg Gly Gln Leu Leu Ala Thr Ala Gly Pro Leu Arg Arg Trp Pro Ala

 50 55 60
 - Pro Ser Pro Ala Ser Ser Pro Ala Pro Gly Ala Ala Ser Ala His 65 70 75 80

	Sei	r Va.	l Gli	n Gly	7 Se: 85	r Ala	a Thi	r Ala	a Gly	90 90	/ Ala	a Arg	g Pro	Gly	95	J Arg
	Pro	Tr	o Gly	y Ala 100	a Arg	g Pro	o Met	Glı	1 Sei 105		/ Let	ı Leı	ı Arç	9 Pro		Pro
5	Val	Sei	Gl:	ı Val	Ile	e Val	l Leu	1 His		Asr	Tyr	Thi	Gl _y	v Lys	Lev	Arg
	Gly	Ala 130	Sei	туг	Glr	Pro	Gly 135		Gly	/ Leu	Arg	Ala 140		Ala	Val	. Val
10	Cys 145	Leu	.Ala	val	Cys	150		Ile	Val	. Leu	. Glu 155		Leu	Ala	Val	Leu 160
5^	Leu	. Val	Leu	Gly	Arg 165		Pro	Arg	Phe	His 170		Pro	Met	Phe	Leu 175	
	Leu	Gly	Ser	Leu 180	Thr	Leu	Ser	Asp	Leu 185		Ala	Gly	Ala	Ala 190	Tyr	Ala
15	Ala	Asn	11e 195		Leu	Ser	Gly	Pro 200	Leu	Thr	Leu	Lys	Leu 205	Ser	Pro	Ala
	Leu	Trp 210	. Phe	Ala	Arg	Glu	Gly 215	Gly	Val	Phe	Val	Ala 220	Leu	Thr	AÌa	Ser
20	Val 225	Leu	Ser	Leu	Leu	Ala 230	Ile	Ala	Leu	Glu	Arg 235	Ser	Leu	Thr	Met	Ala 240
	Arg	Arg	Gly	Pro	Ala 245	Pro	Val	Ser	Ser	Arg 250	Gly	Arg	Thr	Leu	Ala 255	Met
	Ala	Ala	Ala	Ala 260	Trp	Gly	Val	Ser	Leu 265	Leu	Leu	Gly	Leu	Leu 270	Pro	Ala
25	Leu	Gly	Trp 275	Asn	Cys	Leu	Gly	Arg 280	Leu	Asp	Ala	Cys	Ser 285	Thr	Val	Leu
	Pro	Leu 290	Tyr	Ala	Lys	Ala	Tyr 295	Val	Leu	Phe		Val 300	Leu	Ala	Phe	Val
30	Gly 305	Ile	Leu	Ala	Ala	Ile 310	Cys	Ala	Leu	Tyr	Ala 315	Arg	Ile	Tyr	Cys	Gln 320
	Val	Arg	Ala	Asn	Äla 325	Arg	Arg	Leu	Pro	Ala 330	Arg	Pro	Gly	Thr	Ala 335	Gly
	Thr	Thr	Ser	Thr 340	Arg	Ala	Arg	Arg	Lys 345	Pro	Arg	Ser	Leu	Ala 350	Leu	Leu
35	Arg:	Thr	Leu 355	Ser	Val	Val		Leu 360	Ala	Phe	Val.		Сув 365	Trp	Gly	Pro
	Leu	Phe	Leu	Leu	Leu	Leu	Leu	Asp	Val	Ala	Cys .	Pro	Ala	Arg	Thr	Cys

		370					375					380				
	Pro 385	Val	Leu	Leu	Gln	Ala 390	Asp	Pro	Phe	Leu	Gly 395		Ala	Met	Ala	As:
. 5	Ser	Leu	Leu	Asn	Pro 405	Ile	Ile	Tyr	Thr	Leu 410		Asn	Arg	Asp	Leu 415	Arg
	His	Ala	Leu	Leu 420	Arg	Leu	Val	Сув	Cys 425	Gly	Arg	His	Ser	Cys 430	Gly	Arc
	Asp	Pro	Ser 435	Gly	Ser	Gln	Gln	ser 440	Ala	Ser	Ala	Ala	Glu 445		Ser	Gly
0	Gly	Leu 450	Arg	Arg	Cys	Leu	Pro 455	Pro	Gly	Leu	Asp	Gly 460	Ser	Phe	Ser	Gly
	Ser 465	Glu	Arg	Ser	Ser	Pro 470	Gln	Arg	Asp	Gly	Leu 475	Asp	Thr	Ser	Gly	Ser 480
5	Thr	Gly	Ser	Pro	Gly 485	Ala	Pro	Thr	Ala	Ala 490	Arg	Thr	Leu	Val	Ser 495	Glu
	Dro	73-	717.	7												-

- (32) INFORMATION FOR SEQ ID NO:31:
 - (i) SEQUENCE CHARACTERISTICS:

20 (A) LENGTH: 1029

- (A) LENGTH: 1029 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- 25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:31:

ATGCAAGCCG TCGACAATCT CACCTCTGCG CCTGGGAACA CCAGTCTGTG CACCAGAGAC 60
TACAAAATCA CCCAGGTCCT CTTCCCACTG TCCTACACTG TCCTGTTTT TGTTGGACTT 120
ATCACAAATG GCCTGGCGAT GAGGATTTTC TTTCAAAATCC GGAGTAAATC AAACTTTATT 180
ATTTTTCTTA AGAACACAGT CATTTCTGAT CTTCCATGA TTCTGACTTT TCCATTCAAA 240

30 ATTCTTAGTG ATGCCAAACT GGGAACAGGA CCACTGAGAA CTTTTGTGTG TCAAGTTACC 300
TCCGTCATAT TTTATTCAC AATGTATATC AGTATTCAT TCCTGGGACT GATAACTATC 360
GATCGCTACC AGAAGACCAC CAGGCCATTT AAAACATCCA ACCCCAAAAA TCTCTTGGGG 420
GCTAAGATTC TCTCTGTTGT CATCTGGGCA TTCATGTTCT TACTCTCTTT GCCTAACATG 480

ATTCTGACCA ACAGGCAGCC GAGAGACAAG AATGTGAAGA AATGCTCTTT CCTTAAATCA 540
GAGTTCGGTC TAGTCTGGCA TGAAATAGTA AATTACATCT GTCAAGTCAT TTTCTGGATT 600
AATTTCTTAA TTGTTATTGT ATGTTATACA CTCATTACAA AAGAACTGTA CCGGTCATAC 660
GTAAGAACGA GGGGTGTAGG TAAAGTCCCC AGGAAAAAGG TGAACGTCAA AGTTTTCATT 720
5 ATCATTGCTG TATTCTTTAT TTGTTTTGTT CCTTTCCATT TTGCCCGAAT TCCTTACACC 780
CTGAGCCAAA CCCGGGATGT CTTTGACTGC ACTGCTGAAA ATACTCTGTT CTATGTGAAA 840
GAGAGCACTC TGTGGTTAAC TTCCTTAAAT GCATGCCTGG ATCCGTTCAT CTATTTTTC 900
CTTTGCAAGT CCTTCAGAAA TTCCTTGATA AGTATGCTGA AGTGCCCCAA TTCTGCAACA 960
TCTCTGTCCC AGGACAATAG GAAAAAAGAA CAGGATGGTG GTGACCCAAA TGAAGAGACT1020
10 CCAATGTAA

- (33) INFORMATION FOR SEQ ID NO:32:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 342 amino acids
 - (B) TYPE: amino acid
- 15
- (C) STRANDEDNESS:
- (D) TOPOLOGY: not relevant
- (ii) MOLECULE TYPE: protein
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:32:
- Met Gln Ala Val Asp Asn Leu Thr Ser Ala Pro Gly Asn Thr Ser Leu 20 1 5 10, 15
 - Cys Thr Arg Asp Tyr Lys Ile Thr Gln Val Leu Phe Pro Leu Leu Tyr
 20 25 30
 - Thr Val Leu Phe Phe Val Gly Leu Ile Thr Asn Gly Leu Ala Met Arg
- 25 Ile Phe Phe Gln Ile Arg Ser Lys Ser Asn Phe Ile Ile Phe Leu Lys 50 55 60
 - Asn Thr Val Ile Ser Asp Leu Leu Met Ile Leu Thr Phe Pro Phe Lys 65 70 75 80
- Ile Leu Ser Asp Ala Lys Leu Gly Thr Gly Pro Leu Arg Thr Phe Val 85 90 95
 - Cys Gln Val Thr Ser Val Ile Phe Tyr Phe Thr Met Tyr Ile Ser Ile 100 105 110
 - Ser Phe Leu Gly Leu Ile Thr Ile Asp Arg Tyr Gln Lys Thr Thr Arg

			11!	5				120)				125	5		
	Pro) Phe	Lys	s Thi	Ser	Asn	135		Asn	Lei	Leu	140		Lys	Ile	Le
5 .	Ser 145	Val	. Val	L Il∈	: Trp	Ala 150	. Phe	: Met	Phe	Leu	Leu 155		Leu	Pro •) Asn	Ме 16
	Ile	. Leu	Thr	Asr	Arg 165	Gln	Pro	Arg	Asp	Lys 170		Val	Lys	Lys	Cys 175	
	Phe	Leu	Lys	Ser 180	Glu	Phe	Gly	Leu	Val 185		His	Glu	Iļe	Val	Asn	Ту
10	Ile	Суз	Gln 195	Val	Ile	Phe	Trp	Ile 200	Asn	Phe	Leu	Ile	Val 205		Val	Cy:
	Tyr	Thr 210	Leu	Ile	Thr	Lys	Glu 215	Leu	Tyr	Arg	Ser	Tyr 220	Val	Arg	Thr	Arg
15	Gly 225	Val	Gly	Lys	Val	Pro 230	Arg	Lys	Lys	Val	Asn 235	Val	Lys	Val	Phe	I16
	Ile	Ile	Ala	Val	Phe 245	Phe	Ile	Cys	Phe	Val 250	Pro	Phe	His	Phe	Ala 255	Arg
	Ile	Pro	Тут	Thr 260	Leu	Ser	Gln	Thr	Arg 265	Asp	Val	Phe	Asp	Cys 270	Thr	Ala
20	Glu	Asn	Thr 275	Leu	Phe	Tyr	Val	Lys 280	Glu	Ser	Thr	Leu.	Trp 285	Leu	Thr	Ser
	Leu	Asn 290	Ala	Cys	Leu	Asp	Pro 295	Phe	Ile	Tyr		Phe 300	Leu	Cys	Lys	Ser
25	Phe 305	Arg	Asn	Ser	Leu	Ile 310	Ser	Met	Leu	Lys	Cys 315	Pro	Asn	Ser	Ala	Thr 320
	Ser	Leu	Ser	Gln	Asp 325	Asn	Arg	Lys		Glu 330	Gln	Asp	Gly	Gly	Asp 335	Pro
	Asn	Glu	Glu	Thr 340	Pro	Mết										
30 (34)	INFO	RMAT	ІОИ	FOR	SEQ	ID N	0:33	:								
	(i)	(A) (B)	LEN TYP	GTH: E: n	RACT: 107	7 ba:	se p	airs			•					5.
35		(C) (D)			DNES: Y: 1:		-	e ·			• •				v	

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:

- ATGTCGGTCT GCTACCGTCC CCCAGGGAAC GAGACACTGC TGAGCTGGAA GACTTCGCGG 60 GCCACAGGCA CAGCCTTCCT GCTGCTGGCG GCGCTGCTGG GGCTGCCTGG CAACGGCTTC 120 GTGGTGTGGA GCTTGGCGGC CTGGCGGCCT GCACGGGGC GACCGCTGGC GGCCACGCTT 180 5 GTGCTGCACC TGGCGCTGGC CGACGGCGCG GTGCTGCTGC TCACGCCGCT CTTTGTGGCC 240 TTCCTGACCC GGCAGGCCTG GCCGCTGGGC CAGGCGGGCT GCAAGGCGGT GTACTACGTG 300 TGCGCGCTCA GCATGTACGC CAGCGTGCTG CTCACCGGCC TGCTCAGCCT GCAGCGCTGC 360 CTCGCAGTCA CCCGCCCCTT CCTGGCGCCT CGGCTGCGCA GCCCGGCCCT GGCCCGCCGC 420 CTGCTGCTGG CGGTCTGGCT GGCCGCCCTG TTGCTCGCCG TCCCGGCCGC CGTCTACCGC 480 10 CACCTGTGGA GGGACCGCGT ATGCCAGCTG TGCCACCCGT CGCCGGTCCA CGCCGCCGCC 540 CACCTGAGCC TGGAGACTCT GACCGCTTTC GTGCTTCCTT TCGGGCTGAT GCTCGGCTGC 600 TACAGCGTGA CGCTGGCACG GCTGCGGGGC GCCCGCTGGG GCTCCGGGCG GCACGGGGCG 660 CGGGTGGGCC GGCTGGTGAG CGCCATCGTG CTTGCCTTCG GCTTGCTCTG GGCCCCCTAC 720 CACGCAGTCA ACCTTCTGCA GGCGGTCGCA GCGCTGGCTC CACCGGAAGG GGCCTTGGCG 780 15 AAGCTGGGCG GAGCCGGCCA GGCGGCGCA GCGGGAACTA CGGCCTTGGC CTTCTTCAGT 840 TCTAGCGTCA ACCCGGTGCT CTACGTCTTC ACCGCTGGAG ATCTGCTGCC CCGGGCAGGT 900 CCCCGTTTCC TCACGCGGCT CTTCGAAGGC TCTGGGGAGG CCCGAGGGGG CGGCCGCTCT 960 AGGGAAGGGA CCATGGAGCT CCGAACTACC CCTCAGCTGA AAGTGGTGGG GCAGGGCCGC1020 GGCAATGGAG ACCCGGGGGG TGGGATGGAG AAGGACGGTC CGGAATGGGA CCTTTGA 20 (35) INFORMATION FOR SEQ ID NO:34:

 - (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 358 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS:
 - (D) TOPOLOGY: not relevant
 - (ii) MOLECULE TYPE: protein

- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:34:
- Met Ser Val Cys Tyr Arg Pro Pro Gly Asn Glu Thr Leu Leu Ser Trp 1 5 10 15
- 30 Lys Thr Ser Arg Ala Thr Gly Thr Ala Phe Leu Leu Leu Ala Ala Leu

				20					25					30		
	Leu	Gly	7 Let 35	ı Pro	Gly	/ Ası	ı Gl	y Pho	e Va.	l Val	l Trp	Ser	Le:	ı Ala	Gly	7 Trp
5	Arg	Pro 50	Ala	Arg	g Gly	/ Arg	Pro 55	o Lei	ı Ala	a Ala	a Thr	Leu 60	Va]	L Leu	His	Leu
	Ala 65	Leu	Ala	Asp	o Gly	70	a Val	L Let	ı Leı	ı Leı	ı Thr 75	Pro	Leu	1 Phe	Val	Ala 80
	Phe	Leu	Thr	Arg	Gln 85	.Ala	Trp	o Pro	Lev	90	Gln	Ala	, Gly	Cys	Lys 95	Ala
10	Val	Tyr	Tyr	Val 100	Cys	Ala	Leu	. Ser	Met 105		Ala	Ser	Val	Leu 110		. Thr
	Gly	Leu	Leu 115	Ser	Leu	Gln	Arg	Cys 120		Ala	. Val	Thr	Arg 125		Phe	Leu
15	Ala	Pro 130	Arg	Leu	Arg	Ser	Pro 135		Leu	Ala	Arg	Arg		Leu	Leu	Ala
	Val 145	Trp	Leu	Ala	Ala	Leu 150		Leu	Ala	Val	Pro 155	Ala	Ala	Val	Tyr	Arg 160
	His	Leu	Trp	Arg	Asp 165	Arg	Val	Cys	Gln	Leu 170	Cys	His	Pro	Ser	Pro 175	
20	His	Ala	Ala	Ala 180	His	Leu	Ser	Leu	Glu 185	Thr	Leu	Thr	Ala	Phe 190	Val	Leu
	Pro	Phe	Gly 195	Leu	Met	.Leu	Gly	Cys 200	Tyr	Ser	Val	Thr	Leu 205	Ala	Arg	Leu
25	Arg	Gly 210	Ala	Arg	Trp	Gly	Ser 215	Gly	Arg	His	Gly	Ala 220	Arg	Val	Gly	Arg
	Leu 225	Val	Ser	Ala	Ile	Val 230	Leu	Ala	Phe	Gly	Leu 235	Leu	Trp	Ala	Pro	Tyr 240
•	His	Ala	Val	Asn	Leu 245	Leu	Gln	Ala	Val	Ala 250	Ala	Leu	Ala		Pro 255	Glu
30	Gly .	Ala	Leu	Ala 260	Lys .	Leu	Gly	Gly	Ala 265	Gly	Gln	Ala	Ala	Arg 270	Ala	Gly
	Thr	Thr .	Ala 275	Leu	Ala	Phe	Phe	Ser 280	Ser	Ser	Val .		Pro 285	Val :	Leu	Tyr
35	Val 1	Phe '	Thr	Ala	Gly :		Leu 295	Leu	Pro	Arg		Gly :	Pro	Arg :	Phe	Leu
	Thr 2	arg 1	Leu	Phe		Gly 310	Ser	Gly	Glu		Arg (Gly (Gly.	Gly I		Ser 320

- Arg Glu Gly Thr Met Glu Leu Arg Thr Thr Pro Gln Leu Lys Val Val 325 330 335
- Gly Gln Gly Arg Gly Asn Gly Asp Pro Gly Gly Gly Met Glu Lys Asp 340 345 350
- 5 Gly Pro Glu Trp Asp Leu 355
 - (36) INFORMATION FOR SEQ ID NO:35:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1005 base pairs
- 10 (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA (genomic)
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:

AGCAGATGGG CTCATGAACT CCTACTTTCA TTCAGAGAAA AGTGA

15 ATGCTGGGGA TCATGGCATG GAATGCAACT TGCAAAAACT GGCTGGCAGC AGAGGCTGCC CTGGAAAAGT ACTACCTTTC CATTTTTTAT GGGATTGAGT TCGTTGTGGG AGTCCTTGGA 120 AATACCATTG TTGTTTACGG CTACATCTTC TCTCTGAAGA ACTGGAACAG CAGTAATATT 180 TATCTCTTTA ACCTCTCTGT CTCTGACTTA GCTTTTCTGT GCACCCTCCC CATGCTGATA 240 AGGAGTTATG CCAATGGAAA CTGGATATAT GGAGACGTGC TCTGCATAAG CAACCGATAT 300 20 GTGCTTCATG CCAACCTCTA TACCAGCATT CTCTTTCTCA CTTTTATCAG CATAGATCGA 360 TACTTGATAA TTAAGTATCC TTTCCGAGAA CACCTTCTGC AAAAGAAAGA GTTTGCTATT 420 TTAATCTCCT TGGCCATTTG GGTTTTAGTA ACCTTAGAGT TACTACCCAT ACTTCCCCTT 480 ATAAATCCTG TTATAACTGA CAATGGCACC ACCTGTAATG ATTTTGCAAG TTCTGGAGAC 540 CCCAACTACA ACCTCATTTA CAGCATGTGT CTAACACTGT TGGGGTTCCT TATTCCTCTT 600 25 TTTGTGATGT GTTTCTTTTA TTACAAGATT GCTCTCTTCC TAAAGCAGAG GAATAGGCAG 660 GTTGCTACTG CTCTGCCCCT TGAAAAGCCT CTCAACTTGG TCATCATGGC AGTGGTAATC 720 TTCTCTGTGC TTTTTACACC CTATCACGTC ATGCGGAATG TGAGGATCGC TTCACGCCTG 780 GGGAGTTGGA AGCAGTATCA GTGCACTCAG GTCGTCATCA ACTCCTTTTA CATTGTGACA 840 CGGCCTTTGG CCTTTCTGAA CAGTGTCATC AACCCTGTCT TCTATTTTCT TTTGGGAGAT 900 30 CACTTCAGGG ACATGCTGAT GAATCAACTG AGACACAACT TCAAATCCCT TACATCCTTT 960

((37)	INFORMATION	FOR	SEO	TD	NO-36.

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 334 amino acids
 - (B) TYPE: amino acid
- (C) STRANDEDNESS:
 - (D) TOPOLOGY: not relevant
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:
- Met Leu Gly Ile Met Ala Trp Asn Ala Thr Cys Lys Asn Trp Leu Ala

 10 1 5 10 15
 - Ala Glu Ala Ala Leu Glu Lys Tyr Tyr Leu Ser Ile Phe Tyr Gly Ile 20 25 30
 - Glu Phe Val Val Gly Val Leu Gly Asn Thr Ile Val Val Tyr Gly Tyr
 35 40 45
- 15 Ile Phe Ser Leu Lys Asn Trp Asn Ser Ser Asn Ile Tyr Leu Phe Asn 50 55 60
 - Leu Ser Val Ser Asp Leu Ala Phe Leu Cys Thr Leu Pro Met Leu Ile 65 70 75 80
- Arg Ser Tyr Ala Asn Gly Asn Trp Ile Tyr Gly Asp Val Leu Cys Ile 20 85 90 95
 - Ser Asn Arg Tyr Val Leu His Ala Asn Leu Tyr Thr Ser Ile Leu Phe 100 105 110
 - Leu Thr Phe Ile Ser Ile Asp Arg Tyr Leu Ile Ile Lys Tyr Pro Phe 115 120 125
- 25 Arg Glu His Leu Leu Gln Lys Lys Glu Phe Ala Ile Leu Ile Ser Leu 130 135 140
 - Ala Ile Trp Val Leu Val Thr Leu Glu Leu Leu Pro Ile Leu Pro Leu 145 150 155 160
- Ile Asn Pro Val Ile Thr Asp Asn Gly Thr Thr Cys Asn Asp Phe Ala 30 165 170 175
 - Ser Ser Gly Asp Pro Asn Tyr Asn Leu Ile Tyr Ser Met Cys Leu Thr 180 185 190
 - Leu Leu Gly Phe Leu Ile Pro Leu Phe Val Met Cys Phe Phe Tyr Tyr 195 200 205
- Lys Ile Ala Leu Phe Leu Lys Gln Arg Asn Arg Gln Val Ala Thr Ala 210 215 220

315 ...

- Ser Arg Trp Ala His Glu Leu Leu Ser Phe Arg Glu Lys 325 330
- 15 (38) INFORMATION FOR SEQ ID NO:37:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1296 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
- 20 (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA (genomic)
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:
- ATGCAGGCGC TTAACATTAC CCCGGAGCAG TTCTCTCGGC TGCTGCGGGA CCACAACCTG 60

 ACGCGGGAGC AGTTCATCGC TCTGTACCGG CTGCGACCGC TCGTCTACAC CCCAGAGCTG 120

 25 CCGGGACGCG CCAAGCTGGC CCTCGTGCTC ACCGGCGTGC TCATCTTCGC CCTGGCGCTC 180

 TTTGGCAATG CTCTGGTGTT CTACGTGGTG ACCCGCAGCA AGGCCATGCG CACCGTCACC 240

 AACATCTTTA TCTGCTCCTT GGCGCTCAGT GACCTGCTCA TCACCTTCTT CTGCATTCCC 300

 GTCACCATGC TCCAGAACAT TTCCGACAAC TGGCTGGGG GTGCTTTCAT TTGCAAGATG 360

 GTGCCATTG TCCAGTCTAC CGCTGTTGTG ACAGAAATGC TCACTATGAC CTGCATTGCT 420

 30 GTGGAAAGGC ACCAGGGACT TGTGCATCCT TTTAAAATGA AGTGGCAATA CACCAACCGA 480

 AGGGCTTTCA CAATGCTAGG TGTGGTCTGG CTGGTGGCAG TCATCGTAGG ATCACCCATG 540

 TGGCACGTGC AACAACTGA GATCAAAATAT GACTTCCTAT ATGAAAAGGA ACACATCTGC 660

 TGCTTAGAAG AGTGGACCAG CCCTGTGCAC CAGAAGATCT ACACCACCTT CATCCTTGTC 660

ATCCTCTCC TCCTGCCTCT TATGGTGATG CTTATTCTGT ACAGTAAAAT TGGTTATGAA 720
CTTTGGATAA AGAAAAGAGT TGGGGATGGT TCAGTGCTTC GAACTATTCA TGGAAAAGAA 780
ATGTCCAAAA TAGCCAGGAA GAAGAAACGA GCTGTCATTA TGATGGTGAC AGTGGTGGCT 840
CTCTTTGCTG TGTGCTGGGC ACCATTCCAT GTTGTCCATA TGATGATTGA ATACAGTAAT 900
5 TTTGAAAAAGG AATATGATGA TGTCACAATC AAGATGATTT TTGCTATCGT GCAAAATTATT 960
GGATTTTCCA ACTCCATCTG TAATCCCATT GTCTATGCAT TTATGAATGA AAACCTTCAAA1020
AAAAATGTTT TGTCTGCAGT TTGTTATTGC ATAGTAAATA AAACCTTCTC TCCAGCACAA1080
AGGCATGGAA ATTCAGGAAT TACAATGATG CGGAAGAAAG CAAAGTTTTC CCTCAGAGAG1140
AATCCAGTGG AGGAAACCAA AGGAGAAGCA TTCAGTGATG GCAACATTGA AGTCAAATTG1200
10 TGTGAACAGA CAGAGGAGAA GAAAAAGCTC AAACGACATC TTGCTCTCT TAGGTCTGAA1260
CTGGCTGAGA ATTCTCCTTT AGACAGTGGG CATTAA 1296

(39) INFORMATION FOR SEQ ID NO:38:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 431 amino acids
- (B) TYPE: amino acid
 - (C) STRANDEDNESS:
 - (D) TOPOLOGY: not relevant
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:38:
- 20 Met Gln Ala Leu Asn Ile Thr Pro Glu Gln Phe Ser Arg Leu Leu Arg 1 5 10 15
 - Asp His Asn Leu Thr Arg Glu Gln Phe Ile Ala Leu Tyr Arg Leu Arg 20 25 30
- Pro Leu Val Tyr Thr Pro Glu Leu Pro Gly Arg Ala Lys Leu Ala Leu 25 35 40 45
 - Val Leu Thr Gly Val Leu Ile Phe Ala Leu Ala Leu Phe Gly Asn Ala 50 55 60
 - Leu Val Phe Tyr Val Val Thr Arg Ser Lys Ala Met Arg Thr Val Thr 65 70 75 80
- Asn Ile Phe Ile Cys Ser Leu Ala Leu Ser Asp Leu Leu Ile Thr Phe 85 90 95
 - Phe Cys Ile Pro Val Thr Met Leu Gln Asn Ile Ser Asp Asn Trp Leu

Cly Gly Ala Phe Ile Cys Lys Met Val Pro Phe Val Gln Ser Thr Ali 115 120 125 125 125 125 130 130 140 135 140 135 140 140 135 140 140 135 140 140 140 135 140 140 140 140 140 140 140 140 140 140 140 140 140 140 140 155 160 155 160 155 160 155 160 155 160 155 160 165 170 177 177 177 180 175 160 170 175 160 170 175 170			10	0	•		105	5.				110)	٠.
S	٠	Gly Gl	y Ala Ph 115	e Ile C	ys Lys			. Pro	Phe	Val			Thi	c Ala
Arg Ala Phe Thr Met Leu Gly Val Val Trp Leu Val Ala Val Ile Val 165 Arg Ala Phe Thr Met Leu Gly Val Val Trp Leu Val Ala Val Ile Val 165 10 Gly Ser Pro Met Trp His Val Gln Gln Leu Glu Ile Lys Tyr Asp Phe 180 Leu Tyr Glu Lys Glu His Ile Cys Cys Leu Glu Glu Trp Thr Ser Pro 200 Val His Gln Lys Ile Tyr Thr Thr Phe Ile Leu Val Ile Leu Phe Leu 210 Leu Pro Leu Met Val Met Leu Ile Leu Tyr Ser Lys Ile Gly Tyr Glu 225 Leu Trp Ile Lys Lys Arg Val Gly Asp Gly Ser Val Leu Arg Thr Ile 245 20 His Gly Lys Glu Met Ser Lys Ile Ala Arg Lys Lys Lys Arg Ala Val 260 Tle Met Met Val Thr Val Val Ala Leu Phe Ala Val Cys Trp Ala Pro 275 Phe His Val Val His Met Met Ile Glu Tyr Ser Asn Phe Glu Lys Glu 295 Tyr Asp Asp Val Thr Ile Lys Met Ile Phe Ala Ile Val Gln Ile Ile 305 310 Glu Asn Phe Lys Lys Asn Val Leu Ser Ala Val Cys Tyr Cys Ile Val 340 Asn Lys Thr Phe Ser Pro Ala Gln Arg His Gly Asn Ser Gly Ile Thr 355 Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Glu 315 Glu Tr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu 315 Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Glu 315 Glu Tr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu 315	5 .	Val Va 13	l Thr Gli O	u Met L			Thr	. Cys	Ile			. Glu	Arg	y His
165 170 175 10 Gly Ser Fro Met Trp His Val Gln Gln Leu Glu Ile Lys Tyr Asp Phe 180 180 185 190 Leu Tyr Glu Lys Glu His Ile Cys Cys Leu Glu Glu Trp Thr Ser Pro 205 Val His Gln Lys Ile Tyr Thr Thr Phe Ile Leu Val Ile Leu Phe Leu 210 220 Leu Pro Leu Met Val Met Leu Ile Leu Tyr Ser Lys Ile Gly Tyr Glu 225 230 235 Leu Trp Ile Lys Lys Arg Val Gly Asp Gly Ser Val Leu Arg Thr Ile 245 250 His Gly Lys Glu Met Ser Lys Ile Ala Arg Lys Lys Lys Arg Ala Val 265 275 20 His Gly Lys Glu Met Ser Lys Ile Glu Tyr Ser Asn Phe Glu Lys Glu 275 Phe His Val Val His Met Met Ile Glu Tyr Ser Asn Phe Glu Lys Glu 290 295 300 Tyr Asp Asp Val Thr Ile Lys Met Ile Phe Ala Ile Val Gln Ile Ile 305 310 315 Glu Asn Phe Lys Lys Asn Val Leu Ser Ala Val Cys Tyr Cys Ile Val 345 Asn Lys Thr Phe Ser Pro Ala Gln Arg His Gly Asn Ser Gly Ile Thr 355 360 Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Glu 385 Glu Tr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu 385		Gln Gl 145	y Leu Va	l His P	ro Phe	Lys	Met	Lys			Тут	Thr	Asr	
Leu Tyr Glu Lys Glu His Ile Cys Cys Leu Glu Glu Typ Thr Ser Pro 200 Val His Gln Lys Ile Tyr Thr Thr Phe Ile Leu Val Ile Leu Phe Leu 210 Leu Pro Leu Met Val Met Leu Ile Leu Tyr Ser Lys Ile Gly Tyr Glu 235 Leu Trp Ile Lys Lys Arg Val Gly Asp Gly Ser Val Leu Arg Thr Ile 245 Leu Trp Ile Lys Lys Arg Val Gly Asp Gly Ser Val Leu Arg Thr Ile 245 260 His Gly Lys Glu Met Ser Lys Ile Ala Arg Lys Lys Arg Ala Val 260 Ile Met Met Val Thr Val Val Ala Leu Phe Ala Val Cys Trp Ala Pro 285 Phe His Val Val His Met Met Ile Glu Tyr Ser Asn Phe Glu Lys Glu 295 Phe His Val Val His Met Met Ile Glu Tyr Ser Asn Phe Glu Lys Glu 295 Gly Phe Ser Asn Ser Ile Cys Asn Pro Ile Val Tyr Ala Phe Met Asn 325 Glu Asn Phe Lys Lys Asn Val Leu Ser Ala Val Cys Tyr Cys Ile Val 345 Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Glu 365 Glu Thr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu 326		Arg Ala	a Phe Thi	Met Le	eu Gly	Val	Val	Trp 170	Leu	Val	Ala	Val		
195 200 205 Val His Gln Lys Ile Tyr Thr Thr Phe Ile Leu Val Ile Leu Phe Leu 210 Leu Pro Leu Met Val Met Leu Ile Leu Tyr Ser Lys Ile Gly Tyr Glu 225 Leu Trp Ile Lys Lys Arg Val Gly Asp Gly Ser Val Leu Arg Thr Ile 245 Leu Trp Ile Lys Glu Met Ser Lys Ile Ala Arg Lys Lys Lys Arg Ala Val 260 Ile Met Met Val Thr Val Val Ala Leu Phe Ala Val Cys Trp Ala Pro 285 26 Phe His Val Val His Met Met Ile Glu Tyr Ser Asn Phe Glu Lys Glu 295 Tyr Asp Asp Val Thr Ile Lys Met Ile Phe Ala Ile Val Gln Ile Ile 305 Gly Phe Ser Asn Ser Ile Cys Asn Pro Ile Val Tyr Ala Phe Met Asn 325 Gly Phe Lys Lys Asn Val Leu Ser Ala Val Cys Tyr Cys Ile Val 340 Asn Lys Thr Phe Ser Pro Ala Gln Arg His Gly Asn Ser Gly Ile Thr 355 Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Glu 375 Glu Thr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu 385	10	Gly Se			ls Val	Gln		Leu	Glu	Ile	Lys		Asp	Phe
Leu Pro Leu Met Val Met Leu Ile Leu Tyr Ser Lys Ile Gly Tyr Glu 225 Leu Trp Ile Lys Lys Arg Val Gly Asp Gly Ser Val Leu Arg Thr Ile 245 Leu Trp Ile Lys Lys Arg Val Gly Asp Gly Ser Val Leu Arg Thr Ile 245 20 His Gly Lys Glu Met Ser Lys Ile Ala Arg Lys Lys Lys Arg Ala Val 265 Ile Met Met Val Thr Val Val Ala Leu Phe Ala Val Cys Trp Ala Pro 280 Phe His Val Val His Met Met Ile Glu Tyr Ser Asn Phe Glu Lys Glu 295 Tyr Asp Asp Val Thr Ile Lys Met Ile Phe Ala Ile Val Gln Ile Ile 305 Gly Phe Ser Asn Ser Ile Cys Asn Pro Ile Val Tyr Ala Phe Met Asn 325 Glu Asn Phe Lys Lys Asn Val Leu Ser Ala Val Cys Tyr Cys Ile Val 340 Asn Lys Thr Phe Ser Pro Ala Gln Arg His Gly Asn Ser Gly Ile Thr 355 Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Glu 370 Glu Thr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu		Leu Ty	r Glu Lys 195	Glu Hi	s Ile		Ċys	Leu	Glu	Glu		Thr	Ser	Pro
Leu Trp Ile Lys Lys Arg Val Gly Asp Gly Ser Val Leu Arg Thr Ile 255 20 His Gly Lys Glu Met Ser Lys Ile Ala Arg Lys Lys Lys Arg Ala Val 260 Ile Met Met Val Thr Val Val Ala Leu Phe Ala Val Cys Trp Ala Pro 280 Phe His Val Val His Met Met Ile Glu Tyr Ser Asn Phe Glu Lys Glu 290 Tyr Asp Asp Val Thr Ile Lys Met Ile Phe Ala Ile Val Gln Ile Ile 305 Gly Phe Ser Asn Ser Ile Cys Asn Pro Ile Val Tyr Ala Phe Met Asn 325 Glu Asn Phe Lys Lys Asn Val Leu Ser Ala Val Cys Tyr Cys Ile Val 340 Asn Lys Thr Phe Ser Pro Ala Gln Arg His Gly Asn Ser Gly Ile Thr 355 Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Val Lys Leu Glu 375 Glu Thr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu	15	Val His	s Gln Lys)	lle Ty			Phe	Ile	Leu		Ile	Leu	Phe	Leu
245		Leu Pro 225	Leu Met			Ile	Leu	Tyr		Lys	Ile	Gly	Tyr	
The Met Met Val Thr Val Val Ala Leu Phe Ala Val Cys Trp Ala Pro 285 Phe His Val Val His Met Met 11e Glu Tyr Ser Asn Phe Glu Lys Glu 290 Tyr Asp Asp Val Thr Ile Lys Met Ile Phe Ala Ile Val Gln Ile Ile 305 310 315 320 335 335 300 345 350 350 350 350 350 350 350 350 350 350 350 355 360 365 365 365 365 365 365 365 365 360 365		Leu Trp	lle Lys	Lys Ar 245	g Val	Gly	Asp		Ser	Val	Leu	Arg		Ile
275 280 285 Phe His Val Val His Met Met Ile Glu Tyr Ser Asn Phe Glu Lys Glu 290 Tyr Asp Asp Val Thr Ile Lys Met Ile Phe Ala Ile Val Gln Ile Ile 305 Gly Phe Ser Asn Ser Ile Cys Asn Pro Ile Val Tyr Ala Phe Met Asn 325 Glu Asn Phe Lys Lys Asn Val Leu Ser Ala Val Cys Tyr Cys Ile Val 340 Asn Lys Thr Phe Ser Pro Ala Gln Arg His Gly Asn Ser Gly Ile Thr 355 Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Glu 370 Glu Thr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu	20	His Gly	Lys Glu 260	Met Se	r Lys	Ile		Arg	Lys	Lys	Lys		Ala	Val.
Tyr Asp Asp Val Thr Ile Lys Met Ile Phe Ala Ile Val Gln Ile Ile 305 Gly Phe Ser Asn Ser Ile Cys Asn Pro Ile Val Tyr Ala Phe Met Asn 325 Glu Asn Phe Lys Lys Asn Val Leu Ser Ala Val Cys Tyr Cys Ile Val 340 Asn Lys Thr Phe Ser Pro Ala Gln Arg His Gly Asn Ser Gly Ile Thr 355 Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Glu 370 Glu Thr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu 385		Ile Met	Met Val 275	Thr Va	l Val		Leu	Phe	Ala	Val		Trp	Ala	Pro
Gly Phe Ser Asn Ser Ile Cys Asn Pro Ile Val Tyr Ala Phe Met Asn 325 Glu Asn Phe Lys Lys Asn Val Leu Ser Ala Val Cys Tyr Cys Ile Val 340 Asn Lys Thr Phe Ser Pro Ala Gln Arg His Gly Asn Ser Gly Ile Thr 355 Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Glu 370 Glu Thr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu 385	25	Phe His 290	Val Val	His Me		Ile	Glu	Tyr			Phe	Glu	Lys	Glu
325 330 335 Glu Asn Phe Lys Lys Asn Val Leu Ser Ala Val Cys Tyr Cys Ile Val 340 Asn Lys Thr Phe Ser Pro Ala Gln Arg His Gly Asn Ser Gly Ile Thr 355 Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Glu 370 Glu Thr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu 385		Tyr Asp	Asp Val				Ile			Ile	Val	Gln	Ile	
Asn Lys Thr Phe Ser Pro Ala Gln Arg His Gly Asn Ser Gly Ile Thr 355 Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Glu 370 Glu Thr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu 385		Gly Phe	Ser Asn	Ser Ile	e Cys	Asn			Val	Tyr	Ala			Asn
Met Met Arg Lys Lys Ala Lys Phe Ser Leu Arg Glu Asn Pro Val Glu 370 375 380 Glu Thr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu	30	Glu Asn	Phe Lys 340	Lys Ası	ı Val	Leu	Ser :	Ala	Val (Cys			Ile	Val
Glu Thr Lys Gly Glu Ala Phe Ser Asp Gly Asn Ile Glu Val Lys Leu	•	Asn Lys	Thr Phe 355	Ser Pro			Arg 1	His (Gly A			Gly	Ile	Thr.
185	35	Met Met 370	Arg Lys			Phe s	Ser 1	Leu A			Asn :	Pro '	Val	Glu
		Glu Thr 385	Lys Cly	Glu Ala 390	Phe	Ser A	Asp (lle (3lu '	Val 1		

Cys Glu Gln Thr Glu Glu Lys Lys Leu Lys Arg His Leu Ala Leu

Phe Arg Ser Glu Leu Ala Glu Asn Ser Pro Leu Asp Ser Gly His 425

- 5 (40) INFORMATION FOR SEQ ID NO:39:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 24 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single

10

- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:39:

CTGTGTACAG CAGTTCGCAG AGTG

- (41) INFORMATION FOR SEQ ID NO:40:
- 15 (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 24 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- 20 (ii) MOLECULE TYPE: DNA (genomic)
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:40:

GAGTGCCAGG CAGAGCAGGT AGAC

- (42) INFORMATION FOR SEQ ID NO:41:
- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 31 base pairs
 - (B) TYPE: nucleic acid (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

 - (ii) MOLECULE TYPE: DNA (genomic)
- 30 (iv) ANTI-SENSE: NO
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:41:

CCCGAATTCC TGCTTGCTCC CAGCTTGGCC C

(43) INFORMATION FOR SEQ ID NO:42:

					•	
5	(i)	SEQUENCE CHARACTERISTICS: (A) LENGTH: 32 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear				
	(ii)	MOLECULE TYPE: DNA (genomic)			•	
	(iv)	ANTI-SENSE: YES				
	(xi)	SEQUENCE DESCRIPTION: SEQ ID	NO:42:			
TGT	GGATC	CT GCTGTCAAAG GTCCCATTCC GG				32
10 (44)	INF	ORMATION FOR SEQ ID NO:43:				
15	(i)	SEQUENCE CHARACTERISTICS: (A) LENGTH: 20 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	f			
	(ii)	MOLECULE TYPE: DNA (genomic)				,
	(iv)	ANTI-SENSE: NO	-			
	(xi)	SEQUENCE DESCRIPTION: SEQ ID	NO:43:		-	
TCAC	AATG	CT AGGTGTGGTC				20
20 (45)	INF	ORMATION FOR SEQ ID NO:44:		e e e		
	(i)	SEQUENCE CHARACTERISTICS: (A) LENGTH: 22 base pairs (B) TYPE: nucleic acid	· ·			•
25		<pre>(C) STRANDEDNESS: single (D) TOPOLOGY: linear</pre>				
	(ii)	MOLECULE TYPE: DNA (genomic)	- 1			
	(iv)	ANTI-SENSE: YES				
	(xi)	SEQUENCE DESCRIPTION: SEQ ID	NO:44:			

30 (46) INFORMATION FOR SEQ ID NO:45:

TGCATAGACA ATGGGATTAC AG

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 511 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single

(D)	TOPOLOGY:	linear
12	, roponogi:	TTHEGI

- (ii) MOLECULE TYPE: DNA (genomic)
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:45:
- TCACAATGCT AGGTGGGTC TGGCTGGTGG CAGTCATCGT AGGATCACCC ATGTGGCACG 60

 5 TGCAACAACT TGAGATCAAA TATGACTTCC TATATGAAAA GGAACACATC TGCTGCTTAG 120

 AAGAGTGGAC CAGCCCTGTG CACCAGAAGA TCTACACCAC CTTCATCCTT GTCATCCTCT 180

 TCCTCCTGCC TCTTATGGTG ATGCTTATTC TGTACGTAAA ATTGGTTATG AACTTTGGAT 240

 AAAGAAAAGA GTTGGGGATG GTTCAGTGCT TCGAACTATT CATGGAAAAG AAATGTCCAA 300

 AATAGCCAGG AAGAAGAAAC GAGCTGTCAT TATGATGGTG ACAGTGGTGG CTCTCTTTGC 360

 10 TGTGTGCTGG GCACCATTCC ATGTTGCCA TATGATGATT GAATACAGTA ATTTTGAAAA 420

 GGAATATGAT GATGTCACAA TCAAGATGAT TTTTGCTATC GTGCAAATTA TTGGATTTTC 480
 - CAACTCCATC TGTAATCCCA TTGTCTATGC A (47) INFORMATION FOR SEQ ID NO:46:
 - (i) SEQUENCE CHARACTERISTICS:

25

- (A) LENGTH: 21 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- 20 (iv) ANTI-SENSE: NO
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:46:

CTGCTTAGAA GAGTGGACCA G

21

- (48) INFORMATION FOR SEQ ID NO:47:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 22 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA (genomic)
- 30 (iv) ANTI-SENSE: NO

	÷ 5.	Z -	• •		
. (xi)	SEQUENCE DESCRIPTION: SEQ	ID NO:47:	٠,		•
CTGTGCAC	CCA GAAGATCTAC AC				2
.(49) INE	FORMATION FOR SEQ ID NO:48:	•			
5	SEQUENCE CHARACTERISTICS: (A) LENGTH: 21 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear			•.	
(ii)	MOLECULE TYPE: DNA (genomic	c)			
	ANTI-SENSE: YES SEQUENCE DESCRIPTION: SEQ 1	ID NO.48			
•	AA GGTGGTGTAG A		•		
·	ORMATION FOR SEQ ID NO:49:				21
(i) 15	SEQUENCE CHARACTERISTICS: (A) LENGTH: 23 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear				
(ii)	MOLECULE TYPE: DNA (genomic)			
20 , (iv)	ANTI-SENSE: YES				-
(xi)	SEQUENCE DESCRIPTION: SEQ I	D NO:49:		÷ .	
GTGTAGATC	TT TCTGGTGCAC AGG		*	•	23
• • • • • • • • • • • • • • • • • • • •	SEQUENCE CHARACTERISTICS: (A) LENGTH: 21 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear				
(ii) i	MOLECULE TYPE: DNA (genomic)				
) (xi)	SEQUENCE DESCRIPTION: SEQ ID	NO:50:	· •		
GCAATGCAG	G TCATAGTGAG C		·		21
(52) INFOI	RMATION FOR SEQ ID NO:51:	· .		•	

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 22 base pairs

5	(A) LENGTH: 27 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear		
(ii)	MOLECULE TYPE: DNA (genomic	•	
(iii)	HYPOTHETICAL: YES	·	
(iv)	ANTI-SENSE: YES		
(xi)	SEQUENCE DESCRIPTION: SEQ II	NO:51:	
10 TGGAGCAT	GG TGACGGGAAT GCAGAAG		27
(53) INF	ORMATION FOR SEQ ID NO:52:		
(i)	SEQUENCE CHARACTERISTICS: (A) LENGTH: 27 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear		
(ii)	MOLECULE TYPE: DNA (genomic)		
(iv)	ANTI-SENSE: YES		• •
(xi)	SEQUENCE DESCRIPTION: SEQ ID	NO:52:	
20 GTGATGAGO	A GGTCACTGAG CGCCAAG		27
(54) INFO	RMATION FOR SEQ ID NO:53:		
25 .	SEQUENCE CHARACTERISTICS: (A) LENGTH: 23 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear		
(ii)	MOLECULE TYPE: DNA (genomic)		
(iv)	ANTI-SENSE: NO		
,	SEQUENCE DESCRIPTION: SEQ ID	NO:53:	,
30 GCAATGCAGO	G CGCTTAACAT TAC	2	23
(55) INFOR	RMATION FOR SEQ ID NO:54:		
(i) s	FOITENCE CUADA COMPATANTAS		

		(B) TYPE: nucleic acid(C) STRANDEDNESS: single(D) TOPOLOGY: linear		
		(ii) MOLECULE TYPE: DNA (genomic)		
	5	(iv) ANTI-SENSE: YES	•	. •
		(xi) SEQUENCE DESCRIPTION: SEQ ID NO:54:		
	TTG	GGTTACA ATCTGAAGGG CA		2
	(56) INFORMATION FOR SEQ ID NO:55:		
	10	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 23 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 		
•	15	(ii) MOLECULE TYPE: DNA (genomic) (iv) ANTI-SENSE: NO		
•		(xi) SEQUENCE DESCRIPTION: SEQ ID NO:55:		÷ ÷.
	ACT	CCGTGTC CAGCAGGACT CTG		23
	(57)	INFORMATION FOR SEQ ID NO:56:		
	20	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 24 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 		
		(ii) MOLECULE TYPE: DNA (genomic)		
	25	(iv) ANTI-SENSE: YES		
		(xi) SEQUENCE DESCRIPTION: SEQ ID NO:56:		
	TGCG	TGTTCC TGGACCCTCA CGTG		24
	(58)	INFORMATION FOR SEQ ID NO:57:		
	30	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 29 base pairs		

(B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 15 (60)

30

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 27 base pairs (B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: YES

- 5:	5 -
(ii) MOLECULE TYPE: DNA (genomi	c)
(iv) ANTI-SENSE: NO	
(xi) SEQUENCE DESCRIPTION: SEQ	ID NO:57:
CAGGCCTTGG ATTTTAATGT CAGGGATGG	•
5 (59) INFORMATION FOR SEQ ID NO:58:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 27 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: DNA (genomic)
(iv) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ IN	D NO:58:
GGAGAGTCAG CTCTGAAAGA ATTCAGG	
15 (60) INFORMATION FOR SEQ ID NO:59:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 27 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: DNA (genomic)	
(iv) ANTI-SENSE: NO	
(xi) SEQUENCE DESCRIPTION: SEQ ID	NO:59:
TGATGTGATG CCAGATACTA ATAGCAC	2
25 (61) INFORMATION FOR SEQ ID NO:60:	· ·

20

25

30

27

21

CCTGATTCAT TTAGGTGAGA TTGAGAC	
(62) INFORMATION FOR SEQ ID NO:61:	٠.
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 22 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: DNA (genomic)	
10 (iv) ANTI-SENSE: NO	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:61:	
GACAGGTACC TTGCCATCAA G	
(63) INFORMATION FOR SEQ ID NO:62:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 22 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: DNA (genomic)	•
20 (iv) ANTI-SENSE: YES	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:62:	
CTGCACAATG CCAGTGATAA GG	•
(64) INFORMATION FOR SEQ ID NO:63:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 27 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:63:

(iv) ANTI-SENSE: NO

CTGACTTCTT GTTCCTGGCA GCAGCGG

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:60:

(65)	INFORMATION	FOR	SEO	ID	NO - 64 -

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 27 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (iv) ANTI-SENSE: YES
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:64:

10 AGACCAGCCA GGGCACGCTG AAGAGTG

27

- (66) INFORMATION FOR SEQ ID NO:65:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 32 base pairs
 - (B) TYPE: nucleic acid
- 15 (0

25

- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (iv) ANTI-SENSE: NO
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:65:

20 GATCAAGCTT CCATCCTACT GAAACCATGG TC

32

- (67) INFORMATION FOR SEQ ID NO:66:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 35 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA (genomic)
 - (iv) ANTI-SENSE: YES
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:66:

30 GATCAGATCT CAGTTCCAAT ATTCACACCA CCGTC

- (68) INFORMATION FOR SEQ ID NO:67:
 - (i) SEQUENCE CHARACTERISTICS:

		(A) LENGTH: 22 base pa(B) TYPE: nucleic acid(C) STRANDEDNESS: sing(D) TOPOLOGY: linear	•		
5	(ii)	MOLECULE TYPE: DNA (gen	omic)		•
	(iv)	ANTI-SENSE: NO		•	
			4		
-	(xi)	SEQUENCE DESCRIPTION: SI	EQ ID NO:67:		
CTG	GTGTG	CT CCATGGCATC CC			22
(69) INF	CORMATION FOR SEQ ID NO:68	3:		
10	(i)	SEQUENCE CHARACTERISTICS (A) LENGTH: 22 base pai (B) TYPE: nucleic acid (C) STRANDEDNESS: singl (D) TOPOLOGY: linear	lrs		
15	(ii)	MOLECULE TYPE: DNA (geno	omic)		
	(iv)	ANTI-SENSE: YES	٠.		
	(xi)	SEQUENCE DESCRIPTION: SE	Q ID NO:68:		
GTA	AGCCT	CC CAGAACGAGA GG	•		22
·· (70)) INF	ORMATION FOR SEQ ID NO:69	•		
20	(i)	SEQUENCE CHARACTERISTICS (A) LENGTH: 24 base pai (B) TYPE: nucleic acid (C) STRANDEDNESS: singl (D) TOPOLOGY: linear	rs		
25	(ii)	MOLECULE TYPE: DNA (geno	mic)		
	(iv)	ANTI-SENSE: NO			:
	(ind)	CROWNING PROGRAMM			
	(X1)	SEQUENCE DESCRIPTION: SE	Q ID NO:69:		
CAGO	GCAGG	GG TGAAGCCTGA GAGC	. •		24
(71)	INFO	DRMATION FOR SEQ ID NO:70	: .		
30	(i)	SEQUENCE CHARACTERISTICS (A) LENGTH: 24 base pair (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	rs ·		

- (ii) MOLECULE TYPE: DNA (genomic)
- (iv) ANTI-SENSE: YES
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:70:

GGCACCTGCT GTGACCTGTG CAGG

24

- 5 (72) INFORMATION FOR SEQ ID NO:71:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 22 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single

10

- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (iv) ANTI-SENSE: NO
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:71:

GTCCTGCCAC TTCGAGACAT GG

22

- 15 (73) INFORMATION FOR SEQ ID NO:72:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 23 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single

20

- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (iv) ANTI-SENSE: YES
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:72:

GAAACTTCTC TGCCCTTACC GTC

2:

- 25 (74) INFORMATION FOR SEQ ID NO:73:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 26 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single

- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (iv) ANTI-SENSE: NO

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:73:

CCAACACCAG CATCCATGGC ATCAAG

(75) INFORMATION FOR SEQ ID NO:74:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 27 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- 10 (iv) ANTI-SENSE: YES
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:74: GGAGAGTCAG CTCTGAAAGA ATTCAGG

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60/157,281	1 October 1999 (01.10.99)	US
60/157,293	1 October 1999 (01.10.99)	US
60/157,282	1 October 1999 (01.10.99)	US
09/417,044	12 October 1999 (12.10.99)	US
09/416,760	12 October 1999 (12.10.99)	US-

- (72) Inventors; and (75) Inventors/Applicants (for US only): CHEN, Ruoping [CN/US]; DANG, Huong, T. [US/US]; 5352 Oak Park Drive, San Diego, CA 92105 (US). LIAW, Chen, W. [US/US]; 7668 Salix Place, San Diego, CA 92129 (US). LIN, I-Lin [-/US]; 8291-7 Gold Coast Drive, San Diego, CA 92126 (US).
- (74) Agents: MILLER, Suzanne, E. et al.; Woodcock Washburn Kurtz Mackiewicz & Norris LLP, 46th floor, One Liberty Place, Philadelphia, PA 19103 (US).
- (81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP. KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

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(54) Title: HUMAN ORPHAN G PROTEIN-COUPLED RECEPTORS

(57) Abstract

The invention disclosed in this patent document relates to transmembrane receptors, more particularly to endogenous, human orphan G protein-coupled receptors

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		1	101/03 33,	/ 2300/
A. CLASSII IPC 7	FICATION OF SUBJECT MATTER C12N15/12 C07K14/72			
According to	o International Patent Classification (IPC) or to both national classifi	cation and IPC	a e	
	SEARCHED			
Minimum do IPC 7	ocumentation searched (classification system followed by classification ${\tt C12N-C07K}$			
	tion searched other than minimum documentation to the extent that			
Electronic d	ata base consulted during the international search (name of data b	ase and, where practical	, search terms used)
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	nts. 105786 - 107045	•	•	
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X Funt	her documents are listed in the continuation of box C.	X Patent family	members are listed	in annex.
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	ent defining the general state of the art which is not lered to be of particular relevance	or priority date and cited to understan	d not in conflict with	the application but a eary underlying the
9	document but published on or after the international	invention "X" document of particular	lar relevance; the c	laimed invention
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Date of the	actual completion of the international search	Date of mailing of	the international sea	arch report
1	4 July 2000	0 2. 88. 0	0	· ·
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	European Patent Office, P.B. 5818 Patentiaan 2 NL 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Mand1,	В	
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Inte onal Application No PCT/US 99/23687

		PCT/US 99	7 23007
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Ε	WO 00 12707 A (MILLENNIUM PHARM INC) 9 March 2000 (2000-03-09) the whole document		49–52		
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Inte onal Application No PCT/US 99/23687

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national application No. PCT/US 99/23687

Box i Obs	vations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)
,	
This Internation	al Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
	Nos.: e they relate to subject matter not required to be searched by this Authority, namely:
•	
2. Claim	Nos.:
becau an ex	e they relate to parts of the International Application that do not comply with the prescribed requirements to such ent that no meaningful International Search can be carried out, specifically:
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, Douce	e droy the depondent duality and are not during in accordance marking second and third sentences of right buffaj.
Box II Obse	vations where unity of invention is lacking (Continuation of item 2 of first sheet)
This Internation	Il Searching Authority found multiple inventions in this international application, as follows:
see	additional sheet
1. 🔻 As all	equired additional search fees were timely paid by the applicant, this International Search Report covers all
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2. As all of any	earchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment additional fee.
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3. As on	some of the required additional search fees were timely paid by the applicant, this International Search Report
Cover	only those claims for which fees were paid, specifically claims Nos.:
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4. No recreated	uired additional search fees were timely paid by the applicant. Consequently, this International Search Report is ad to the invention first mentioned in the claims; it is covered by claims Nos.:
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Remark on Pro	est The additional search fees were accompanied by the applicant's protest.
	No protest accompanied the payment of additional search fees.

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This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-4

Human G protein-coupled receptor as characterized by SEQ.ID.2, a cDNA encoding said receptor as characterized by SEQ.ID.1, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

2. Claims: 5-8

Human G protein-coupled receptor as characterized by SEQ.ID.4, a cDNA encoding said receptor as characterized by SEQ.ID.3, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

3. Claims: 9-12

Human G protein-coupled receptor as characterized by SEQ.ID.6, a cDNA encoding said receptor as characterized by SEQ.ID.5, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

4. Claims: 13-16

Human G protein—coupled receptor as characterized by SEQ.ID.8, a cDNA encoding said receptor as characterized by SEQ.ID.7, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

5. Claims: 17-20

Human G protein-coupled receptor as characterized by SEQ.ID.10, a cDNA encoding said receptor as characterized by SEQ.ID.9, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

6. Claims: 21-24

Human G protein-coupled receptor as characterized by SEQ.ID.12, a cDNA encoding said receptor as characterized by SEQ.ID.11, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

7. Claims: 25-28

Human G protein-coupled receptor as characterized by SEQ.ID.14, a cDNA encoding said receptor as characterized by

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SEQ.ID.13, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

8. Claims: 29-32

Human G protein-coupled receptor as characterized by SEQ.ID.16, a cDNA encoding said receptor as characterized by SEQ.ID.15, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

9. Claims: 33-36

Human G protein-coupled receptor as characterized by SEQ.ID.18, a cDNA encoding said receptor as characterized by SEQ.ID.17, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

10. Claims: 37-40

Human G protein-coupled receptor as characterized by SEQ.ID.20, a cDNA encoding said receptor as characterized by SEQ.ID.19, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

11. Claims: 41-44

Human G protein-coupled receptor as characterized by SEQ.ID.22, a cDNA encoding said receptor as characterized by SEQ.ID.21, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

12. Claims: 45-48

Human G protein-coupled receptor as characterized by SEQ.ID.24, a cDNA encoding said receptor as characterized by SEQ.ID.23, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

13. Claims: 49-52

Human G protein-coupled receptor as characterized by SEQ.ID.26, a cDNA encoding said receptor as characterized by SEQ.ID.25, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

14. Claims: 53-56

Human G protein-coupled receptor as characterized by

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

SEQ.ID.28, a cDNA encoding said receptor as characterized by SEQ.ID.27, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

15. Claims: 57-60

Human G protein-coupled receptor as characterized by SEQ.ID.30, a cDNA encoding said receptor as characterized by SEQ.ID.29, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

16. Claims: 61-64

Human G protein-coupled receptor as characterized by SEQ.ID.32, a cDNA encoding said receptor as characterized by SEQ.ID.31, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

17. Claims: 65-68

Human G protein-coupled receptor as characterized by SEQ.ID.34, a cDNA encoding said receptor as characterized by SEQ.ID.33, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

18. Claims: 69-72

Human G protein-coupled receptor as characterized by SEQ.ID.36, a cDNA encoding said receptor as characterized by SEQ.ID.35, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

19. Claims: 73-76

Human G protein-coupled receptor as characterized by SEQ.ID.38, a cDNA encoding said receptor as characterized by SEQ.ID.37, a plasmid comprising said cDNA, and a host cell comprising said plasmid.

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